

**TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371**

10/088376

ATTORNEY DOCKET NUMBER IN-5562	U.S. APPLICATION NO. (IF KNOWN SEE 37 CFR 15)	
INTERNATIONAL APPLICATION NO. PCT/EP00/08503	INTERNATIONAL FILING DATE 31 August 2000 (31.08.2000)	PRIORITY DATE CLAIMED 23 September 1999 (23.09.1999)
TITLE OF INVENTION: USE OF ASSOCIATIVE THICKENERS BASED ON POLYURETHANE AND/OR DIPROPYLENE GLYCOL MONOALKYL ETHERS IN COLOURED AND/OR DECORATIVE EFFECT MULTI-LAYERED LACQUERS		

APPLICATION(S) FOR DO/EO/US: **Ines MANGELS, Karl-Heinz JOSTEN, Udo NENNER, Egon WEGNER, Ekkehard SAPPER and Nicola ZARSE**

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

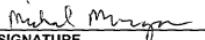
1. This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.
2. This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.
3. This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371 (b) and PCT Articles 22 and 39(1).
4. A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date
5. A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. have been transmitted by the International Bureau.
 - c. is not required, as the application was filed in the United States Receiving Office (RO/US)
6. A translation of the International Application into English (35 U.S.C. 371(C)(2)).
7. Amendment to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. have been transmitted by the International Bureau.
 - c. have not been made; however, the time limit for making such amendments has NOT expired.
 - d. have not been made and will not be made
8. A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. A translation of the annex to the International Preliminary Examination Report under PCT Article 36

Items 11. to 16. below concern other document(s) or information included:

11. An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included
13. A FIRST preliminary amendment.
 A SECOND or SUBSEQUENT preliminary amendment.
14. A substitute specification.
15. A Change of power of attorney and/or address letter.
16. Other items or information:
A copy of the cover sheet from the PCT Published Application
A copy of the cover sheet from the Priority Document

I hereby certify that the attached correspondence is being deposited with the United States Postal Service in an envelope as "Express Mail Post Office to Addressee" Mailing Label No. **E7973 679 288US** addressed to the Commissioner for Patents, Washington, DC 20231 on March 11, 2002.

Marjorie E. Ellis
Marjorie Ellis

U.S. APPLICATION NO. (If known) 10/008576 INTERNATIONAL APPLICATION NO. PCT/EP00/08503		ATTORNEY'S DOCKET NUMBER IN-5562
17. <input checked="" type="checkbox"/> The following fees are submitted:		
Basic National Fee (37 CFR 1.492(a)(1)-(5)):		
Neither international preliminary examination fee (37 CFR 1.482) Nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO..... \$970.00		
International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO..... \$890.00		
International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.44(a)(2)) paid to USPTO..... \$690.00		
International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4)..... \$670.00		
International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4) \$ 96.00		
ENTER APPROPRIATE BASIC FEE AMOUNT = \$890.00		
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)). \$		
Claims	Number Filed	Number Extra
Total Claims	18 - 20 = 0	X \$ 18.00 \$
Independent claims	02 - 03 = 0	X \$ 84.00 \$
Multiple dependent claim(s) (if applicable)		+ \$280.00 \$
TOTAL OF ABOVE CALCULATION = \$890.00		
Reduction by 1/2 for filing by small entity, if applicable. Verified Small Entity statement must also be filed. (Note 37 CFR 1.9, 1.27, 1.28). \$		
SUBTOTAL = \$890.00		
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)). + \$		
TOTAL NATIONAL FEE = \$890.00		
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property + \$		
TOTAL FEES ENCLOSED = \$930.00		
<input type="checkbox"/> Amount to be: \$ <input type="checkbox"/> refunded <input type="checkbox"/> Charged \$930.00		
<p>a. <input type="checkbox"/> A check in the amount of \$_____ to cover the above fees is enclosed.</p> <p>b. <input checked="" type="checkbox"/> Please charge my Deposit Account No. <u>23-3425</u> in the amount of <u>\$930.00</u> to cover the above fees A triplicate copy of this sheet is enclosed.</p> <p>c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>23-3425</u>. A triplicate copy of this sheet is enclosed.</p>		
<p>NOTE: Where an appropriate time limit under 37 CFR 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.</p> <p>SEND ALL CORRESPONDENCE TO: BASF CORPORATION Patent Department 26701 Telegraph Road Southfield, Michigan 48034-2442 (248) 948-2355 Customer No. 26922</p>		
 SIGNATURE <hr/> <p>Michael F. Morgan Name _____ 42,906 REGISTRATION NUMBER _____</p>		

10/088376

Patent Application Data Sheet

JC10 Recd PCT/US 11 MAR 2002

Title Line One:: Title Line Two:: Title Line Three:: Title Line Four:: Title Line Five:	USE OF ASSOCIATIVE THICKENERS BASED ON POLYURETHANE AND/OR DIPROPYLENE GLYCOL MONOALKYL ETHERS IN COLOURED AND/OR DECORATIVE EFFECT MULTI-LAYERED LACQUERS
Total Drawing Sheets:: Formal Drawings?::	
Application Type::	Utility
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Representative Information::	BASF Corporation Patent Department 26701 Telegraph Road Southfield, Michigan 48034
Attorney(s) Name:: Registration Number::	Anne Gerry Sabourin 33,772
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Attorney(s) Name:: Registration Number::	Michael F. Morgan 42,906
Representative Customer Number::	26922
Continuity Information This application is a National Phase of >Application One:: Filing Date::	PCT/EP00/08503 31 August 2000
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Prior Foreign Applications Foreign Application One:: Filing Date:: Country:: Priority Claimed::	199 45 574.0 23 September 1999 Germany yes

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PATENT

(Practitioner's Docket No. IN-5562)

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of

Ines MANGELS, Karl-Heinz JOSTEN, Udo
NENNER, Egon WEGNER, Ekkehard
SAPPER and Nicola ZARSE

Serial No.: This application is a National
Phase of Patent Application PCT/EP00/08503
filed 31 August 2000.

Filed: Monday, March 11, 2002

For: USE OF ASSOCIATIVE
THICKENERS BASED ON
POLYURETHANE AND/OR
DIPROPYLENE GLYCOL MONOALKYL
ETHERS IN COLOURED AND/OR
DECORATIVE EFFECT MULTI-LAYERED
LACQUERS

Group Art Unit: Not Assigned

Examiner: Not Assigned

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an envelope as "Express Mail Post Office to Addressee"
Mailing Label No. ET973 679 288US addressed to the
Assistant Commissioner for Patents, Washington, D.C.
20231 on March 11, 2002.

Marjorie Ellis

PRELIMINARY AMENDMENT UNDER 37 CFR § 1.115

Commissioner for Patents

Washington, D.C. 20231

Dear Sir:

This preliminary amendment is submitted with the application for entry into the U.S.
National Phase under Chapter II. This application is based on PCT/EP00/08503 filed on 31
August 2000.

In connection with the filing of this National Phase application, please make the
following preliminary amendments.

IN THE SPECIFICATION

After the title, please insert --This application is a National Phase Application of
Patent Application **PCT/EP00/08503** filed on 31 August 2000.--

IN THE CLAIMS:

Please cancel claims 1-23.

Please add claims 24-41.

24. (New) A method for suppressing optical defects in a paint system, wherein the paint system is one of a multicoat color paint system, a multicoat effect paint system, a multicoat color and effect paint system, a color refinish paint system, an effect refinish paint system, and a color and effect refinish paint system, comprising including an associative thickener in the paint system, wherein the associative thickener is at least one of a polyurethane-based associative thickener and a dipropylene glycol monoalkyl ether.
25. (New) The method of claim 24, wherein the paint system comprises at least one basecoat layer and at least one clearcoat layer.
26. (New) The method of claim 25, wherein the basecoat layer is produced from an aqueous basecoat material.
27. (New) The method of claim 26, wherein the aqueous basecoat material comprises a polyurethane polymer.
28. (New) The method of claim 24, wherein the associative thickener is present in a basecoat layer.
29. (New) The method of claim 24, wherein the polyurethane-based associative thickener and the dipropylene glycol monoalkyl ether are present in a basecoat layer.
30. (New) The method of claim 24, wherein the optical defects comprise at least one of:
 - i) light-colored spots, introduced by at least one of dried-up liquid residues and abrasion dust residues, which remain after abrading of defects in the paint system or in a surferc that is to be coated with the paint system; and
 - ii) polishing spots, induced by the polishing of defects in the paint system, which is performed for the purpose of refinishing the paint system.

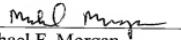
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31. (New) The method of claim 26, wherein the aqueous basecoat material comprises
 - (A) at least one water-soluble or -dispersible polyurethane,
 - (B) at least one crosslinking agent,
 - (C) at least one pigment that is at least one of a color pigment, an effect pigment, and a color and effect pigment,
 - (D) at least one neutralizing agent,
 - (E) at least one inorganic thickener,
 - (F) an associative thickener that is at least one of a dipropylene glycol monoalkyl ether and a polyurethane-based associative thickener,
 - (G) optionally, at least one water-soluble or -dispersible polyacrylate resin prepared in the presence of a water-soluble or -dispersible polyurethane, and
 - (H) optionally, at least one water-soluble or -dispersible polyester resin.
 32. (New) The method of claim 31, wherein the aqueous basecoat material, based on its overall weight, contains from 0.5 to 11% by weight of the dipropylene glycol monoalkyl ether.
 33. (New) The method of claim 31, wherein the aqueous basecoat material, based on its overall weight, contains from 0.1 to 4% by weight of the polyurethane-based associative thickener.
 34. (New) The method of claim 31, wherein the aqueous basecoat material further comprises at least one additive.
 35. (New) The paint system prepared by the process of claim 24.
 36. (New) The paint system of claim 35, wherein the paint system is one of an original coating for a motor vehicle body, an industrial coating, an electrical components coating, a coil coating, a packaging coating, a plastics coating, and a furniture coating.

37. (New) An aqueous basecoat material comprising
- (A) at least one water-soluble or -dispersible polyurethane,
 - (B) at least one crosslinking agent,
 - (C) at least one pigment that is at least one of a color pigment, an effect pigment, and a color and effect pigment,
 - (D) at least one neutralizing agent,
 - (E) at least one inorganic thickener,
 - (F) an associative thickener that is at least one of a dipropylene glycol monoalkyl ether and a polyurethane-based associative thickener,
 - (G) optionally, at least one water-soluble or -dispersible polyacrylate resin prepared in the presence of a water-soluble or -dispersible polyurethane, and
 - (H) optionally, at least one water-soluble or -dispersible polyester resin.
38. (New) The aqueous basecoat material of claim 37, wherein the aqueous basecoat material, based on its overall weight, contains from 0.5 to 11% by weight of the dipropylene glycol monoalkyl ether.
39. (New) The aqueous basecoat material of claim 37, wherein the aqueous basecoat material, based on its overall weight, contains from 0.1 to 4% by weight of the polyurethane-based associative thickener.
40. (New) The aqueous basecoat material of claim 37 further comprises at least one additive.
41. (New) A paint system comprising at least one basecoat layer; wherein the paint system is one of a multicoat color paint system, a multicoat effect paint system, a multicoat color and effect paint system, a color refinish paint system, an effect refinish paint system, and a color and effect refinish paint system; and wherein the basecoat layer is produced from the aqueous basecoat material of claim 37.

REMARKS

Upon entry of the present amendment, claims 24-41 are pending in the application. New claims 24-41 add no new matter, as these claims contain subject matter deleted from claims 1-23. There are two (2) independent claims and a total of eighteen (18) claims pending in the application. Applicants respectfully request entry of the preliminary amendment.

Respectfully Submitted,


Michael F. Morgan
Registration No. 42,906
Customer No. 26922

Date: March 11, 2002
BASF Corporation
26701 Telegraph Road
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USE OF POLYURETHANE-BASED ASSOCIATIVE THICKENERS AND/OR
DIPROPYLENE GLYCOL MONOALKYL ETHERS IN MULTICOAT COLOR
AND/OR EFFECT PAINT SYSTEMS

5 The present invention relates to the use of polyurethane-based associative thickeners and/or dipropylene glycol monoalkyl ethers to suppress optical defects in multicoat color and/or effect paint systems or their refinishes. The present invention also relates
10 to novel aqueous basecoat materials which comprise at least one polyurethane-based associative thickener and/or at least one dipropylene glycol monoalkyl ether. The present invention further relates to the use of the novel aqueous basecoat materials for producing
15 multicoat color and/or effect paint systems and their refinishes.

Owing to their numerous advantageous properties, multicoat color and/or effect paint systems are
20 nowadays employed in many fields, examples being the coating of motor vehicle bodies, industrial components, including electrical components, coils, and packaging, and fabrics or furniture. Owing to the particularly stringent requirements on the part of automakers and
25 their customers, these multicoat color and/or effect paint systems have become established in particular in this field of use.

As is known, the bodies of automobiles are nowadays coated with a multicoat paint system composed of a primer, particularly an electrocoat, a surfacer or an antistonechip primer, and a multicoat color and/or effect paint system. The multicoat paint system itself is generally composed of at least one basecoat and at least one clearcoat. Advantageously, the basecoat is produced from an aqueous basecoat material. In accordance with the particularly advantageous wet-on-wet technique, the aqueous basecoat material is applied to the surfacer or the antistonechip primer and partly dried, but not crosslinked. Thereafter, at least one clearcoat film is applied to the uncured aqueous basecoat film and is cured together with the aqueous basecoat film. This technique is also employed for the refinish of the multicoat color and/or effect paint systems.

Both during the production of the multicoat color and/or effect paint systems and during the refinish of defects present therein, optical defects can occur which are manifested to disruptive effect and which give cause for complaints. These optical defects may have a variety of causes.

25

Accordingly, defects in the surfacer or in the antistonechip primer are normally removed by abrading. When this is done, abrasion dust residues in the form of fingerprints or, in the case of wet abrasion, in the

form of dried water droplets can remain. After overcoating with aqueous basecoat and clearcoat material, these "fingerprints" and/or "droplets" show up as light-colored, clearly visible sites in the 5 resulting multicoat color and/or effect paint system. These optical defects are also referred to by the skilled worker as "light spots".

Optical defects of this kind also arise if the 10 clearcoat of a multicoat color and/or effect paint system is abraded for purposes of better adhesion of a refinish that is to be applied. These optical defects are also referred to by the skilled worker as "abrasion spots".

15

These pale spots are not abrasion scars which are poorly covered, but may instead be induced even by beads of sweat or the sweat on the fingers.

20 Another kind of optical defect, known as polishing spots, results if the finished multicoat color and/or effect paint system is polished at damaged sites, but then, owing to further damaged sites, has to be subjected to a complete refinish, with the polished 25 sites as well being overcoated. In the refinish, these polishing sites are very clearly visible owing to their different shade.

To date it has been possible to prevent these optical defects only by extremely careful and cautious working and a high additional cleaning effort, since the aqueous basecoat materials available at present do not
5 have the sufficient insensitivity to such defects in and on the surfacers and/or in and on the multicoat color and/or effect paint systems, and are therefore unable, or unable sufficiently, to compensate these defects optically. This fact, however, leads to
10 relatively long processing times and thus to higher costs in line finishing or in line or workshop refinish.

There is therefore a high demand for basecoat
15 materials, especially aqueous basecoat materials, which no longer have these disadvantages but instead provide optical compensation of the defects while retaining all of their other advantageous properties, if not indeed improving upon them.

20 Aqueous basecoat materials which are outstandingly suitable for producing color and/or effect paint systems are known, for example, from the German patent application DE-A-44 37 535. They comprise an inorganic
25 thickener, such as a phyllosilicate, and organic solvents, such as butyl glycol. As binders they comprise water-soluble or -dispersible polyurethanes, water-soluble or -dispersible polyacrylate resins prepared in the presence of such polyurethanes, and

water-soluble or -dispersible polyesters. They have a very high storage stability even at relatively high temperatures. The use of polyurethane-based associative thickeners and/or dipropylene glycol monoalkyl ethers
5 is not disclosed by DE-A-44 37 535.

Polyurethane-based associative thickeners and their performance properties and advantages are known. By way of example, reference may be made here to Römpf Lexikon
10 Lacke und Druckfarben, Georg Thieme Verlag, Stuttgart, New York, 1998, "thickeners", pages 599 to 600, and the textbook "Lackadditive" [Additives for Coatings] by Johan Bieleman, Wiley-VCH, Weinheim, New York, 1998, pages 51 to 59 and 65. The use of these polyurethane-
15 based associative thickeners for suppressing optical defects in multicoat color and/or effect paint systems and their refinishes are not described therein.

It is an object of the present invention to provide
20 novel basecoat materials, especially the aqueous basecoat materials, which comprise novel combinations of constituents and which allow the production and refinish of multicoat color and/or effect paint systems without the formation of light spots or polishing
25 spots. The novel basecoat materials, particularly the novel aqueous basecoat materials, ought to continue to have the particular advantages of the existing basecoat materials or to exhibit additional, novel advantageous properties.

Found accordingly was the novel use of polyurethane-based associative thickeners and/or dipropylene glycol monoalkyl ethers for suppressing optical defects in
5 multicoat color and/or effect paint systems and their refinishes.

In the text below, this novel use is referred to as "inventive use".

10

Additionally found was the novel aqueous basecoat material which comprises

(A) at least one water-soluble or -dispersible
15 polyurethane,

(B) at least one crosslinking agent,

(C) at least one color and/or effect pigment,

20

(D) at least one neutralizing agent,

(E) at least one inorganic thickener, and

25 (F) at least one dipropylene glycol monoalkyl ether.

Additionally found was the further novel aqueous basecoat material which comprises

(A) at least one water-soluble or -dispersible polyurethane,

(B) at least one crosslinking agent,

5

(C) at least one color and/or effect pigment,

(D) at least one neutralizing agent,

10 (E) at least one inorganic thickener,

(G) at least one polyurethane-based associative thickener,

15 (H) if desired, at least one water-soluble or -dispersible polyacrylate resin prepared in the presence of a water-soluble or -dispersible polyurethane, and

20 (I) at least one water-soluble or -dispersible polyester resin.

In the text below, these novel aqueous basecoat materials are referred to collectively as "aqueous

25 basecoat materials of the invention".

Found not least were the novel multicoat color and/or effect paint systems which comprises at least one

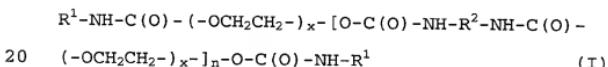
basecoat producible from one of the aqueous basecoat materials of the invention.

In the light of the prior art it was surprising and
5 unforeseeable for the skilled worker that the object of the present invention could be achieved by means of the inventive use. Even more surprising was that the achievement of the object was demonstrably not based on the rheological properties of the aqueous basecoat
10 materials of the invention, which in this respect are equivalent to the known aqueous basecoat materials. Moreover, it could not at all have been expected that the durable suppression or prevention of polishing spots could be achieved by the combination of the
15 polyurethane-based associative thickener (G) for inventive use and at least one dipropylene glycol monoalkyl ether (F).

In accordance with the invention, the polyurethane-based associative thickener (G) or the dipropylene glycol monoalkyl ether (F) can be employed alone, with the use alone of the associative thickener (G) offering particular advantages and therefore being preferred in accordance with the invention. Very particular
25 advantages result, however, from the joint use of the two constituents, which is therefore particularly preferred in accordance with the invention.

The first inventively essential constituent of the aqueous basecoat materials of the invention is therefore at least one polyurethane-based associative thickener (G). These thickeners are normally composed
5 of nonionic hydrophobic polymers which are available either in liquid form, as a 50 percent strength solution in water or in organic solvents, for example, or in powder form. They are based on water-soluble polyurethanes having a comparatively low molecular
10 weight of from 10 000 to 50 000.

The water-soluble polyurethanes (G) are prepared by reacting diisocyanates, especially the diisocyanates described below, with diols, especially the diols
15 described below, and hydrophobic blocking components, thereby resulting in the following idealized structure of the general formula I.



In the general formula I, the indices x and n stand for integers whose value is sufficient to give the molecular weights indicated above.

25

The radical R¹ stands for a hydrophobic aliphatic or aromatic group. Examples of suitable groups of this kind are oleyl, stearyl, dodecylphenyl or nonylphenyl.

The radical R² stands for a hydrophilic segment. Examples of suitable hydrophilic segments are polyesters, especially polyesters of maleic acid and ethylene glycol, and polyethers, especially 5 polyethylene glycol or its derivatives.

The polyurethane-based associative thickeners for inventive use therefore contain the three following kinds of segments or building-block groups:

10

- hydrophobic end segments,
- two or more hydrophilic segments, and
- 15 - urethane groups.

These segments or building-block groups can be linked to one another in a very wide variety of ways, so resulting in a very wide variety of polymer structures.

20

In accordance with the invention, however, advantageous associative thickeners are those which contain linear and comb-shaped polymer structures. It is essential that each polymer molecule contains at least two hydrophobic end segments.

25

For the preparation of the aqueous basecoat materials of the invention, the associative thickeners (G) for inventive use are employed as powders or, preferably,

as 20 to 30% strength by weight solutions in at least one water-dilutable solvent.

In the aqueous basecoat materials of the invention they
5 are advantageously present in an amount of from 0.1 to 4.0%, preferably from 0.4 to 3.5%, and in particular from 0.4 to 3.0% by weight, based in each case on the overall weight of the respective aqueous basecoat material.

10

Dipropylene glycol monoalkyl ethers are compounds which are known per se and are employed in the coatings field as solvents or as solvent additions. Examples of suitable alkyl radicals in this context are methyl, propyl, isopropyl, n-butyl, n-pentyl or n-hexyl, of which the methyl radical is particularly advantageous and is therefore used with particular preference.

The dipropylene glycol methyl ether (F) used with
20 particular preference is a compound known per se or a mixture known per se, both of which are available commercially. It is derived from dipropylene glycol, 1,1'-oxybis-(2-propanol), and/or isomers thereof.

25 In the aqueous basecoat materials of the invention, the dipropylene glycol monoalkyl ethers (F) are present advantageously in an amount of from 0.5 to 11%, preferably from 1.0 to 9%, and in particular from 1.0

to 7.0% by weight, based in each case on the overall weight of the respective aqueous basecoat material.

The other essential constituent of the aqueous basecoat
5 materials of the invention is at least one water-soluble or -dispersible polyurethane (A). Polyurethanes (A) and processes for preparing them are known per se and are described, for example, in the German patent DE-A-44 37 535.

10

Advantageously, depending on the nature of the stabilization, the polyurethane (A) for inventive use has an acid number or amine number of from 10 to 250 mg KOH/g (ionic stabilization or nonionic plus ionic
15 stabilization) or from 0 to 10 mg KOH/g (nonionic stabilization), an OH number of from 30 to 350 mg KOH/g, and a number-average molecular weight of from 1 500 to 55 000 daltons.

20 For the preparation of the polyurethanes (A) it is conventional to use diisocyanates and also, if desired and in minor amounts, polyisocyanates, for the purpose of introducing branches. In the context of the present invention, minor amounts are amounts which do not cause
25 gelling of the polyurethanes (A) during their preparation. This may also be prevented by using small amounts of monoisocyanates as well.

Examples of suitable diisocyanates are isophorone diisocyanate (i.e., 5-isocyanato-1-isocyanatomethyl-1,3,3-trimethylcyclohexane), 5-isocyanato-1-(2-isocyanatoeth-1-yl)-1,3,3-trimethylcyclohexane, 5-isocyanato-5 1-(3-isocyanatoprop-1-yl)-1,3,3-trimethylcyclohexane, 5-isocyanato-(4-isocyanatobut-1-yl)-1,3,3-trimethylcyclohexane, 1-isocyanato-2-(3-isocyanatoprop-1-yl)-cyclohexane, 1-isocyanato-2-(3-isocyanatoeth-1-yl)-cyclohexane, 1-isocyanato-2-(4-isocyanatobut-1-yl)-10 cyclohexane, 1,2-diisocyanatocyclobutane, 1,3-diisocyanatocyclobutane, 1,2-diisocyanatocyclopentane, 1,3-diisocyanatocyclopentane, 1,2-diisocyanatocyclohexane, 1,3-diisocyanatocyclohexane, 1,4-diisocyanato-cyclohexane, dicyclohexylmethane 2,4'-diisocyanate, 15 trimethylene diisocyanate, tetramethylene diisocyanate, pentamethylene diisocyanate, hexamethylene diisocyanate, ethylethylene diisocyanate, trimethylhexane diisocyanate, heptanemethylene diisocyanate or diisocyanates derived from dimeric fatty acids, as marketed under the commercial designation DDI 20 1410 by the company Henkel and described in patents DO 97/49745 and WO 97/49747, especially 2-heptyl-3,4-bis(9-isocyanatononyl)-1-pentylcyclohexane, or 1,2-, 1,4- or 1,3-bis(isocyanatomethyl)cyclohexane, 1,2-, 1,4- or 25 1,4- or 1,3-bis(2-isocyanatoeth-1-yl)cyclohexane, 1,3-bis(3-isocyanatoprop-1-yl)cyclohexane, 1,2-, 1,4- or 1,3-bis(4-isocyanatobut-1-yl)cyclohexane, liquid bis(4-isocyanatocyclohexyl)methane with a trans/trans content of up to 30% by weight, preferably 25% by weight, and

in particular 20% by weight, as is described in patents DE-A-44 14 032, GB-A-1 220 717, DE-A-16 18 795 or DE-A-17 93 785; tolylene diisocyanate, xylylene diisocyanate, bisphenylene diisocyanate, naphthylene 5 diisocyanate or diphenylmethane diisocyanate.

Examples of suitable polyisocyanates are the isocyanurates of the diisocyanates described above.

10 Examples of highly suitable monoisocyanates are phenyl isocyanate, cyclohexyl isocyanate or stearyl isocyanate.

15 The polyurethanes (A) for inventive use are also prepared using

20 - saturated and unsaturated polyols of relatively high and low molecular mass, especially diols and, in minor amounts, triols for the purpose of introducing branches,

- polyamines,

- amino alcohols, and

25 - compounds which introduce stabilizing (potentially) ionic and/or nonionic functional groups.

Examples of suitable polyols are saturated or olefinically unsaturated polyester polyols which are prepared by reacting

- 5 - unsulfonated or sulfonated saturated and/or unsaturated polycarboxylic acids or their esterifiable derivatives, alone or together with monocarboxylic acids, and
- 10 - saturated and/or unsaturated polyols, alone or together with monoools.

Examples of suitable polycarboxylic acids are aromatic, aliphatic and cycloaliphatic polycarboxylic acids.

- 15 Preference is given to the use of aromatic and/or aliphatic polycarboxylic acids.

- 20 Examples of suitable aromatic polycarboxylic acids are phthalic acid, isophthalic acid, terephthalic acid, phthalic, isophthalic or terephthalic acid monosulfonate, or halophthalic acids, such as tetrachlorophthalic or tetrabromophthalic acid, among which isophthalic acid is advantageous and is therefore used with preference.

25

Examples of suitable acyclic aliphatic or unsaturated polycarboxylic acids are oxalic acid, malonic acid, succinic acid, glutaric acid, adipic acid, pimelic acid, suberic acid, azelaic acid, sebacic acid,

undecanedicarboxylic acid or dodecanedicarboxylic acid, or dimeric fatty acids, or maleic acid, fumaric acid or itaconic acid, of which adipic acid, glutaric acid, azelaic acid, sebacic acid, dimeric fatty acids and 5 maleic acid are advantageous and are therefore used with preference.

Examples of suitable cycloaliphatic and cyclic unsaturated polycarboxylic acids are 1,2-cyclobutane-
10 dicarboxylic acid, 1,3-cyclobutanedicarboxylic acid, 1,2-cyclopentanedicarboxylic acid, 1,3-cyclopentane-dicarboxylic acid, hexahydrophthalic acid, 1,3-cyclohexanedicarboxylic acid, 1,4-cyclohexanedicarboxylic acid, 4-methylhexahydrophthalic acid, tricyclodecane-
15 dicarboxylic acid, tetrahydrophthalic acid or 4-methyl-tetrahydrophthalic acid. These dicarboxylic acids may be used both in their cis and in their trans form and also as a mixture of both forms.

20 Also suitable are the esterifiable derivatives of the abovementioned polycarboxylic acids, such as their monoesters or polyesters with aliphatic alcohols having 1 to 4 carbon atoms or hydroxy alcohols having 1 to 4 carbon atoms, for example. It is also possible to use
25 the anhydrides of the abovementioned polycarboxylic acids, where they exist.

Together with the polycarboxylic acids it is also possible if desired to use monocarboxylic acids, such

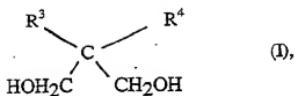
as, for example, benzoic acid, tert-butylbenzoic acid, lauric acid, isononanoic acid, fatty acids of naturally occurring oils, acrylic acid, methacrylic acid, ethacrylic acid or crotonic acid. The preferred 5 monocarboxylic acid used is isononanoic acid.

Examples of suitable polyols are diols and triols, especially diols. Normally, triols are used alongside the diols in minor amounts in order to introduce 10 branches into the polyester polyols. In the context of the present invention, minor amounts are amounts which do not cause gelling of the polyester polyols during their preparation.

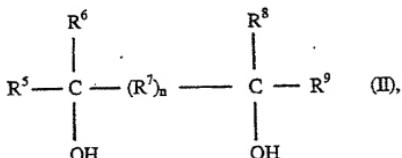
15 Suitable diols are ethylene glycol, 1,2- or 1,3-propanediol, 1,2-, 1,3- or 1,4-butanediol, 1,2-, 1,3-, 1,4- or 1,5-pentanediol, 1,2-, 1,3-, 1,4-, 1,5- or 1,6-hexanediol, neopentyl hydroxypivalate, neopentyl 20 glycol, diethylene glycol, 1,2-, 1,3- or 1,4-cyclohexanediol, 1,2-, 1,3- or 1,4-cyclohexane-dimethanol, trimethylpentanediol, ethylbutylpropanediol or the positionally isomeric diethyloctanediols. These diols may also be used per se for the preparation of the polyurethanes (A) for inventive use.

25

Further examples of suitable diols are diols of the formula I or II:



where R^3 and R^4 are each an identical or different radical and are an alkyl radical having 1 to 18 carbon atoms, an aryl radical or a cycloaliphatic radical, with the proviso that R^3 and/or R^4 must not be methyl;



where R^5 , R^6 , R^8 and R^9 are each identical or different radicals and are an alkyl radical having 1 to 6 carbon atoms, a cycloalkyl radical or an aryl radical and R^7 is an alkyl radical having 1 to 6 carbon atoms, an aryl radical or an unsaturated alkyl radical having 1 to 6 carbon atoms, and n is either 0 or 1.

Suitable diols I of the general formula I are all propanediols of the formula in which either R^3 or R^4 , or R^3 and R^4 , is or are not methyl, such as, for example, 2-butyl-2-ethyl-1,3-propanediol, 2-butyl-2-methyl-1,3-propanediol, 2-phenyl-2-methyl-1,3-propanediol, 2-propyl-2-ethyl-1,3-propanediol, 2-di-tert-butyl-1,3-

propanediol, 2-butyl-2-propyl-1,3-propanediol, 1-di-hydroxymethylbicyclo[2.2.1]heptane, 2,2-diethyl-1,3-propanediol, 2,2-dipropyl-1,3-propanediol or 2-cyclohexyl-2-methyl-1,3-propanediol, et cetera.

5

Examples of diols II of the general formula II that may be used are 2,5-dimethyl-2,5-hexanediol, 2,5-diethyl-2,5-hexanediol, 2-ethyl-5-methyl-2,5-hexanediol, 2,4-dimethyl-2,4-pentanediol, 2,3-dimethyl-2,3-butanediol,

10 1,4-(2'-hydroxypropyl)benzene and 1,3-(2'-hydroxypropyl)benzene.

Of these diols, hexanediol and neopentyl glycol are particularly advantageous and are therefore used with
15 particular preference.

The abovementioned diols may also be used per se to prepare the polyurethanes (A) for inventive use.

20 Examples of suitable triols are trimethylolethane, trimethylolpropane or glycerol, especially trimethylolpropane.

25 The abovementioned triols may also be used per se to prepare the polyurethanes (A) for inventive use (cf. the patent EP-A-0 339 433).

If desired, minor amounts of monoools may also be used. Examples of suitable monoools are alcohols or phenols

such as ethanol, propanol, n-butanol, sec-butanol, tert-butanol, amyl alcohols, hexanols, fatty alcohols, allyl alcohol, or phenol.

- 5 The polyester polyols may be prepared in the presence of small amounts of a suitable solvent as entrainer. Examples of entrainers used are aromatic hydrocarbons, such as especially xylene and (cyclo)aliphatic hydrocarbons, e.g., cyclohexane or methylcyclohexane.

10

Further examples of suitable polyols are polyester diols which are obtained by reacting a lactone with a diol. They are notable for the presence of terminal hydroxyl groups and repeating polyester fractions of

- 15 the formula $\text{-(CO-(CHR}^{10})_m\text{-CH}_2\text{-O-)}$. Here, the index m is preferably from 4 to 6 and the substituent R¹⁰ is hydrogen or an alkyl, cycloalkyl, or alkoxy radical. No substituent contains more than 12 carbon atoms. The total number of carbon atoms in the substituent does
20 not exceed 12 per lactone ring. Examples are hydroxycaproic acid, hydroxybutyric acid, hydroxydecanoic acid, and/or hydroxystearic acid.

Preferred for the preparation of the polyester diols is
25 the unsubstituted # #-caprolactone, where m is 4 and all R¹⁰ substituents are hydrogen. The reaction with lactone is started by low molecular mass polyols such as ethylene glycol, 1,3-propanediol, 1,4-butanediol, or dimethylolcyclohexane. It is also possible, however, to

react other reaction components, such as ethylenediamine, alkylidialkanolamines, or else urea, with caprolactone. Other suitable diols of relatively high molecular mass are polylactam diols, which are 5 prepared by reacting, for example, ###-caprolactam with low molecular mass diols.

Further examples of suitable polyols include polyether polyols, especially those having a number-average 10 molecular weight of from 400 to 5 000, in particular from 400 to 3 000. Examples of highly suitable polyether diols are polyether diols of the general formula H-(-O-(CHR¹¹)_o-)_pOH, where the substituent R¹¹ is hydrogen or a lower, unsubstituted or substituted alkyl 15 radical, the index o is from 2 to 6, preferably from 3 to 4, and the index p is from 2 to 100, preferably from 5 to 50. Especially suitable examples are linear or branched polyether diols such as poly(oxyethylene) glycols, poly(oxypropylene) glycols, and 20 poly(oxybutylene) glycols.

The polyether diols ought on the one hand not to introduce excessive amounts of ether groups, since otherwise the polyurethanes (A) for inventive use that 25 are formed start to swell in water. On the other hand, they can be used in amounts which ensures the nonionic stabilization of the polyurethanes (A). In that case they serve as the functional nonionic groups (a3) described below.

The polyurethane (A) for inventive use comprises alternatively

5 (a1) functional groups convertible into cations by neutralizing agents and/or quaternizing agents, and/or cationic groups, especially ammonium groups,

10 or

(a2) functional groups convertible into anions by neutralizing agents, and/or anionic groups, especially carboxylic acid and/or carboxylate groups,

15

and/or

(a3) nonionic hydrophilic groups, especially 20 poly(alkylene ether) groups.

Examples of suitable functional groups (a1) for inventive use and convertible into cations by neutralizing agents (D) and/or quaternizing agents are 25 primary, secondary or tertiary amino groups, secondary sulfide groups or tertiary phosphine groups, especially tertiary amino groups or secondary sulfide groups.

Examples of suitable cationic groups (a1) for inventive use are primary, secondary, tertiary or quaternary ammonium groups, tertiary sulfonium groups or quaternary phosphonium groups, preferably quaternary 5 ammonium groups or quaternary ammonium groups, tertiary sulfonium groups, but especially tertiary sulfonium groups.

Examples of suitable functional groups (a2) for 10 inventive use and convertible into anions by neutralizing agents (D) are carboxylic acid, sulfonic acid or phosphonic acid groups, especially carboxylic acid groups.

15 Examples of suitable anionic groups (a2) for inventive use are carboxylate, sulfonate or phosphonate groups, especially carboxylate groups.

Examples of suitable neutralizing agents (D) for 20 functional groups (a1) convertible into cations are organic and inorganic acids such as formic acid, acetic acid, lactic acid, dimethylolpropionic acid, citric acid, sulfuric acid, hydrochloric acid or phosphoric acid.

25 Examples of suitable neutralizing agents (D) for functional groups (a2) convertible into anions are tertiary amines, such as trimethylamine, triethylamine, tributylamine, dimethylaniline, diethylaniline,

triphenylamine, dimethylethanolamine, diethylethanolamine, methyldiethanolamine or triethanolamine, for example. Neutralization may take place in organic phase or in aqueous phase. A preferred neutralizing agent 5 used is dimethylethanolamine.

The total amount of neutralizing agent (D) used in the aqueous basecoat material of the invention is chosen so that from 1 to 100 equivalents, preferably from 50 to 10 90 equivalents, of the functional groups (a1) or (a2) of the polyurethane (A) for inventive use are neutralized.

15 Of these functional (potentially) ionic groups (a1) and (a2) and functional nonionic groups (a3), the (potentially) anionic groups (a2) are advantageous and are therefore used with particular preference.

The introduction of (potentially) anionic groups (a2) 20 into the polyurethane molecules takes place by way of the incorporation of compounds which contain in the molecule at least one isocyanate-reactive group and at least one group capable of forming anions; the amount to be used may be calculated from the target acid 25 number.

Examples of suitable compounds of this kind are those containing two isocyanate-reactive groups in the molecule. Suitable isocyanate-reactive groups are, in

- particular, hydroxyl groups, and also primary and/or secondary amino groups. Accordingly it is possible, for example, to use alkanoic acids having two substituents on the ###-positioned carbon atom. The substituent may
- 5 be a hydroxyl group, an alkyl group, or, preferably, an alkylol group. These alkanoic acids have at least one, generally from 1 to 3, carboxyl groups in the molecule. They have 2 to about 25, preferably 3 to 10, carbon atoms. Examples of suitable alkanoic acids are
- 10 dihydroxypropionic acid, dihydroxysuccinic acid, and dihydroxybenzoic acid. A particularly preferred group of alkanoic acids are the ###, ###-dimethylolalkanoic acids of the general formula $R^{12}-C(CH_2OH)_2COOH$, R^{12} being a hydrogen atom or an alkyl group having up to about 20
- 15 carbon atoms. Examples of especially suitable alkanoic acids are 2,2-dimethylolacetic acid, 2,2-dimethylolpropionic acid, 2,2-dimethylolbutyric acid, and 2,2-dimethylolpentanoic acid. The preferred dihydroxyalkanoic acid is 2,2-dimethylolpropionic acid.
- 20 Examples of compounds containing amino groups are ###, ###-diaminovaleric acid, 3,4-diaminobenzoic acid, 2,4-diaminotoluenesulfonic acid, and 2,4-diaminodiphenyl ether sulfonic acid.
- 25 Stabilizing nonionic poly(oxyalkylene) groups (a3) may be introduced as lateral or terminal groups into the polyurethane molecules. For this purpose it is possible to use, for example, alkoxypoly(oxyalkylene) alcohols having the general formula $R^{13}O-(CH_2-CHR^{14}-O)_xH$, where

R¹³ is an alkyl radical having 1 to 6 carbon atoms, R¹⁴ is a hydrogen atom or an alkyl radical having 1 to 6 carbon atoms, and the index r is a number between 20 and 75. (cf. the patents EP-A-0 354 261 or
5 EP-A-0 424 705).

The use of polyols, polyamines and amino alcohols leads to an increase in the molecular weight of the polyurethanes (A).

- 10 Suitable polyols for the chain extension are polyols containing up to 36 carbon atoms per molecule, such as ethylene glycol, diethylene glycol, triethylene glycol, 1,2-propanediol, 1,3-propanediol, 1,4-butanediol, 1,2-
15 butylene glycol, 1,6-hexanediol, trimethylolpropane, castor oil or hydrogenated castor oil, di-trimethylolpropane ether, pentaerythritol, 1,2-cyclohexanediol, 1,4-cyclohexanedimethanol, bisphenol A, bisphenol F, neopentyl glycol, neopentyl glycol
20 hydroxypivalate, hydroxyethylated or hydroxypropylated bisphenol A, hydrogenated bisphenol A or mixtures thereof (cf. patents EP-A-0 339 433, EP-A-0 436 941, EP-A-0 517 707).

25 Examples of suitable polyamines have at least two primary and/or secondary amino groups. Polyamines are essentially alkylene polyamines having 1 to 40 carbon atoms, preferably about 2 to 15 carbon atoms. They may carry substituents which have no hydrogen atoms that

are reactive with isocyanate groups. Examples are polyamines having a linear or branched aliphatic, cycloaliphatic or aromatic structure and at least two primary amino groups.

5

Diamines include hydrazine, ethylenediamine, propylenediamine, 1,4-butylenediamine, piperazine, 1,4-cyclohexyldimethylamine, 1,6-hexamethylenediamine, trimethylhexamethylenediamine, methanediamine, isophoronediamine, 4,4'-diaminodicyclohexylmethane, and aminoethylenothanolamine. Preferred diamines are hydrazine, alkyl- or cycloalkyldiamines such as propylenediamine and 1-amino-3-aminomethyl-3,5,5-trimethylcyclohexane.

15

It is also possible to use polyamines containing more than two amino groups in the molecule. In these cases, however, it should be ensured - for example, by using monoamines as well - that no crosslinked polyurethane resins are obtained. Polyamines of this kind which may be used are diethylenetriamine, triethylenetetramine, dipropylenediamine, and dibutylenetriamine. An example of a monoamine is ethylhexylamine (cf. the patent EP-A-0 089 497).

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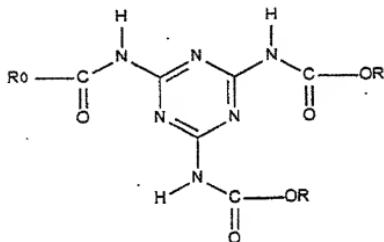
Examples of suitable amino alcohols are ethanolamine or diethanolamine.

- In the aqueous basecoat materials of the invention, the polyurethanes (A) are present advantageously in an amount of from 1.0 to 50%, preferably from 2.0 to 40%, with particular preference from 3.0 to 30%, with very
- 5 particular preference from 4.0 to 25%, and in particular from 5.0 to 20% by weight, based in each case on the overall weight of the respective aqueous basecoat material of the invention.
- 10 The further essential constituent of the coating material of the invention is at least one crosslinking agent (B).
- 15 Examples of suitable crosslinking agents (B) are amino resins, compounds or resins containing anhydride groups, compounds or resins containing epoxide groups, tris(alkoxycarbonylamino)triazines, compounds or resins containing carbonate groups, blocked and/or nonblocked polyisocyanates, beta-hydroxyalkylamides, and compounds
- 20 containing on average at least two groups capable of transesterification, examples being reaction products of malonic diesters and polyisocyanates or of esters and partial esters of polyhydric alcohols of malonic acid with monoisocyanates, as described by the European
- 25 patent EP-A-0 596 460.

Crosslinking agents of this kind are well known to the skilled worker and are offered by numerous companies as sales products.

Examples of suitable polyepoxides are, in particular, all known aliphatic and/or cycloaliphatic and/or aromatic polyepoxides, based for example on bisphenol A or bisphenol F. Examples of suitable polyepoxides also include the polyepoxides available commercially under the designations Epikote® from Shell, Denacol® from Nagase Chemicals Ltd., Japan, such as, for example, Denacol EX-411 (pentaerythritol polyglycidyl ether), Denacol EX-321 (trimethylolpropane polyglycidyl ether), Denacol® EX-512 (polyglycerol polyglycidyl ether) and Denacol® EX-521 (polyglycerol polyglycidyl ether).

The suitable tris(alkoxycarbonylamino)triazines had the following formula:



Examples of suitable tris(alkoxycarbonylamino)triazines are described in the patents US-A-4,939,213, US-A-5,084,541 or EP-A-0 624 577. Use is made in particular of the tris(methoxy-, tris(butoxy- and/or tris(2-ethylhexoxycarbonylamino)triazines.

Of advantage are the methyl butyl mixed esters, the butyl 2-ethylhexyl mixed esters and the butyl esters. These have the advantage over the straight methyl ester 5 of better solubility in polymer melts, and also have less of a tendency to crystallize.

An example of a suitable polyanhydride is polysuccinic anhydride.

10

Examples of suitable beta-hydroxyalkylamides are N,N,N',N'-tetrakis(2-hydroxyethyl)adipamide or N,N,N',N'-tetrakis(2-hydroxypropyl)adipamide.

15 Further examples of suitable crosslinking agents are the blocked polyisocyanates.

Examples of suitable blocking agents are the blocking agents known from the US patent US-A-4,444,954:

20

- i) phenols such as phenol, cresol, xylenol, nitrophenol, chlorophenol, ethylphenol, t-butylphenol, hydroxybenzoic acid, esters of this acid or 2,5-di-tert-butyl-4-hydroxytoluene;
- 25 ii) lactams, such as ϵ -caprolactam, δ -valerolactam, γ -butyrolactam or β -propiolactam;

- iii) active methylenic compounds, such as diethyl malonate, dimethyl malonate, ethyl or methyl acetoacetate or acetylacetone;
- 5 iv) alcohols such as methanol, ethanol, n-propanol, isopropanol, n-butanol, isobutanol, t-butanol, n-amyl alcohol, t-amyl alcohol, lauryl alcohol, ethylene glycol monomethyl ether, ethylene glycol monoethyl ether, ethylene glycol monobutyl ether, diethylene glycol monomethyl ether, diethylene glycol monoethyl ether, propylene glycol monomethyl ether, methoxymethanol, glycolic acid, glycolic esters, lactic acid, lactic esters, 15 methylolurea, methylolmelamine, diacetone alcohol, ethylenechlorohydrin, ethylenebromo-hydrin, 1,3-dichloro-2-propanol, 1,4-cyclo-hexyldimethanol or acetocyanohydrin;
- 20 v) mercaptans such as butyl mercaptan, hexyl mercaptan, t-butyl mercaptan, t-dodecyl mercaptan, 2-mercaptopbenzothiazole, thiophenol, methylthiophenol or ethylthiophenol;
- 25 vi) amides such as acetoanilide, aceto-anisidinamide, acrylamide, methacrylamide, acetamide, stearamide or benzamide;

- vii) imides such as succinimide, phthalimide or maleimide;
- 5 viii) amines such as diphenylamine, phenylnaphthylamine, xylidine, N-phenylxylidine, carbazole, aniline, naphthylamine, butylamine, dibutylamine or butylphenylamine;
- 10 ix) imidazoles such as imidazole or 2-ethyylimidazole;
- 15 x) ureas such as urea, thiourea, ethyleneurea, ethylenethiourea or 1,3-diphenylurea;
- xi) carbamates such as phenyl N-phenylcarbamate or 2-oxazolidone;
- 20 xii) imines such as ethyleneimine;
- xiii) oximes such as acetone oxime, formaldoxime, acetaldoxime, acetoxime, methyl ethyl ketoxime, diisobutyl ketoxime, diacetyl monoxime, benzophenone oxime or chlorohexanone oximes;
- 25 xiv) salts of sulfurous acid such as sodium bisulfite or potassium bisulfite;

xv) hydroxamic esters such as benzyl methacrylohydroxamate (BMH) or allyl methacrylohydroxamate; or

5 xvi) substituted pyrazoles, ketoximes, imidazoles or triazoles; and also

xvii) mixtures of these blocking agents, especially dimethylpyrazole and triazoles, malonates and 10 acetoacetates or dimethylpyrazole and succinimide.

Examples of suitable organic polyisocyanates for blocking are in particular the polyisocyanates known as

15 paint polyisocyanates, containing aliphatically, cycloaliphatically, araliphatically and/or aromatically bonded isocyanate groups. Preference is given to polyisocyanates containing from 2 to 5 isocyanate groups per molecule and having viscosities of from 100 20 to 10 000, preferably from 100 to 5 000. Furthermore, the polyisocyanates can have been hydrophilically or hydrophobically modified in a customary and known way.

Further examples of suitable polyisocyanates for 25 blocking are described in "Methoden der organischen Chemie", Houben-Weyl, Volume 14/2, 4th edition, Georg Thieme Verlag, Stuttgart 1963, page 61 to 70, and by W. Siefken, Liebigs Annalen der Chemie, Volume 562, pages 75 to 136. Suitable examples include the

polyurethane prepolymers containing isocyanate groups, which can be prepared by reacting polyols with an excess of polyisocyanates and which are preferably of low viscosity.

5

Further examples of suitable polyisocyanates for blocking are polyisocyanates containing isocyanurate, biuret, allophanate, iminooxadiazinedione, urethane, urea and/or uretdione groups. Polyisocyanates

10 containing urethane groups, for example, are obtained by reacting some of the isocyanate groups with polyols, such as trimethylolpropane and glycerol, for example. It is preferred to use aliphatic or cycloaliphatic polyisocyanates, especially hexamethylene diisocyanate,

15 dimerized and trimerized hexamethylene diisocyanate, isophorone diisocyanate, dicyclohexylmethane 2,4'-diisocyanate, dicyclohexylmethane 4,4'-diisocyanate, diisocyanates derived from dimeric fatty acids, as marketed under the commercial designation DDI 1410 by

20 the company Henkel and described in the patents DO 97/49745 and WO 97/49747, especially 2-heptyl-3,4-bis(9-isocyanatononyl)-1-pentylcyclohexane; or 1,2-, 1,4- or 1,3-bis(isocyanatomethyl)cyclohexane, 1,2-, 1,4- or 1,3-bis(2-isocyanatoethyl)cyclohexane, 1,3-bis(3-isocyanatoprop-1-yl)cyclohexane or 1,2-, 1,4- or 1,3-bis(4-isocyanatobut-1-yl)cyclohexane, 1,8-diiso-
25 cyanato-4-isocyanatomethyloctane, 1,7-diisocyanato-4-isocyanatomethylheptane or 1-isocyanato-2-(3-iso-

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cyanatopropyl)cyclohexane, or mixtures of these poly-isocyanates.

Very particular preference is given to using mixtures
5 of polyisocyanurates which contain uretdione and/or
isocyanurate groups and/or allophanate groups that are
based on hexamethylene diisocyanate, as formed by
catalytic oligomerization of hexamethylene diisocyanate
using appropriate catalysts.

10

In particular it is possible to use amino resins,
examples being melamine resins, guanamine resins or
urea resins, as crosslinking agents (B). In this case
it is possible to use any amino resin which is suitable
15 for transparent topcoat materials or clearcoat
materials, or a mixture of such amino resins. For
further details refer to Römpf Lexikon Lacke und
Druckfarben, Georg Thieme Verlag, 1998, page 29, "Amino
resins", and the textbook "Lackadditive" by Johan
20 Bieleman, Wiley-VCH, Weinheim, New York, 1998, pages
242 ff., or to the book "Paints, Coatings and
Solvents", second completely revised edition, Editors
D. Stoye and W. Freitag, Wiley-VCH, Weinheim, New York,
1998, pages 80 ff. Also suitable are the customary and
25 known amino resins some of whose methylol and/or
methoxymethyl groups have been defunctionalized by
means of carbamate or allophanate groups. Crosslinking
agents of this kind are described in patents
US-A-4 710 542 and EP-B-0 245 700 and also in the

article by B. Singh and coworkers, "Carbamylmethylated Melamines, Novel Crosslinkers for the Coatings Industry" in Advanced Organic Coatings Science and Technology Series, 1991, Volume 13, pages 193 to 207.

5

Preferably, the amino resins are contained in the aqueous basecoat materials of the invention as the predominant or sole crosslinking agents (B). The other abovementioned crosslinking agents (B) can be used as

10 additional crosslinking agents (B) for the further advantageous variation of the profile of properties of the aqueous basecoat materials of the invention and of the inventive basecoats and inventive multicoat color and/or effect paint systems produced from them, their
15 proportion among the crosslinking agents (B) in that case being < 50% by weight.

In the aqueous basecoat materials of the invention, the crosslinking agents (B) are employed preferably in an amount of from 0.1 to 30%, more preferably from 0.3 to 20%, with particular preference from 0.5 to 10%, and in particular from 1.0 to 8.0% by weight, based in each case on the overall weight of the respective aqueous basecoat material of the invention.

25

Yet another essential constituent of the aqueous basecoat material of the invention is at least one color and/or effect pigment (C). The pigments (C) can be composed of organic or inorganic compounds. The

aqueous basecoat material of the invention therefore ensures, owing to this large number of suitable pigments, a universal breadth of use, and permits the realization of a large number of color shades and
5 optical effects.

As effect pigments (C) it is possible to use metal flake pigments such as commercial aluminum bronzes, aluminum bronzes chromated as per DE-A-36 36 183,
10 commercial stainless steel bronzes, and nonmetallic effect pigments, such as pearlescent pigments and interference pigments, for example. For further details refer to Römpf Lexikon Lacke und Druckfarben, Georg Thieme Verlag, 1998, pages 176, "Effect pigments" and
15 pages 380 and 381, "Metal oxide/mica pigments" to "Metal pigments".

Examples of suitable inorganic color pigments (C) are titanium dioxide, iron oxides, Sicotrans yellow, and
20 carbon black. Examples of suitable organic color pigments (C) are thioindigo pigments indanthrene blue, Cromophthal red, Irgazine orange and Heliogen green. For further details refer to Römpf Lexikon Lacke und Druckfarben, Georg Thieme Verlag, 1998, pages 180 and
25 181, "Iron blue pigments" to "Black iron oxide", pages 451 to 453, "Pigments" to "Pigment volume concentration", page 563 "Thioindigo pigments" and page 567 "Titanium dioxide pigments".

The fraction of the pigments (C) in the aqueous basecoat material of the invention can vary extremely widely and is guided primarily by the opacity of the pigments (C), by the desired shade and by the desired optical effect. In the aqueous basecoat material of the invention, the pigments (C) are present in an amount of preferably from 0.5 to 50%, more preferably from 0.5 to 45%, with particular preference from 0.5 to 40%, with very particular preference from 0.5 to 35%, and in particular from 0.5 to 30% by weight, based in each case on the overall weight of the aqueous basecoat material of the invention. The pigment/binder ratio, i.e., the ratio of the pigments (C) to the above-described polyurethanes (A) or to the polyurethanes (A) and the hereinbelow-described polyacrylate resins (H) and polyesters (I), can also vary extremely widely. This ratio is preferably from 4.0:1.0 to 1.0:50, more preferably from 3.5:1.0 to 1.0:50, with particular preference from 3.0:1.0 to 1.0:40, with very particular preference from 2.5:1.0 to 1.0:30, and in particular from 2.3:1.0 to 1.0:25.

These pigments (C) may also be incorporated into the aqueous basecoat materials of the invention by way of pigment pastes, in which case suitable dispersing resins include the above-described polyurethanes (A) and/or the hereinbelow-described polyacrylate resins (H) and/or polyester resins (I).

Yet another essential constituent of the aqueous basecoat material of the invention is at least one inorganic thickener (E).

- 5 Examples of suitable inorganic thickeners (E) are inorganic phyllosilicates, preferably smectites, especially montmorillonites and hectorites, such as aluminum magnesium silicates, sodium magnesium and sodium magnesium fluorine lithium phyllosilicates of
10 the montmorillonite type or inorganic phyllosilicates such as aluminum magnesium silicates, aluminum magnesium silicates, sodium magnesium and sodium magnesium fluorine lithium phyllosilicates of the montmorillonite type. For further details, refer to the
15 book by Johan Bielemann "Lackadditive", Wiley-VCH, Weinheim, New York, 1998, pages 17 to 30.

In the aqueous basecoat materials of the invention, the thickeners (E) are present preferably in an amount of
20 from 0.01 to 5.0%, more preferably from 0.05 to 3.0%, with particular preference from 0.1 to 2.5%, with very particular preference from 0.2 to 2.0%, and in particular from 0.3 to 1.5% by weight, based in each case on the overall weight of the aqueous basecoat
25 material of the invention.

In one preferred embodiment of the aqueous basecoat material of the invention, it comprises in addition to the polyurethanes (A) at least one water-soluble or

-dispersible polyacrylate resin (H) prepared in the presence of a water-soluble or -dispersible polyurethane.

- 5 Examples of suitable polyurethanes in whose presence the polyacrylate resin (H) is prepared contain olefinically unsaturated groups which are lateral, terminal and/or located within the main polymer chain, or contain no olefinically unsaturated groups. Examples
10 of highly suitable polyurethanes of this kind are known from the patents EP-A-0 521 928, EP-A-0 522 420, EP-A-0 522 419 or EP-A-0 730 613 (olefinically unsaturated) or DE-A-44 37 535 (saturated). Examples of especially suitable polyurethanes are the above-
15 described polyurethanes (A), especially the polyurethanes (A) containing carboxylic acid groups and/or carboxylate groups.

Accordingly, the polyacrylate resins (H) which contain
20 carboxylic acid groups and/or carboxylate groups as solubilizing or dispersing groups are of advantage in accordance with the invention and are therefore used with preference. As regards the neutralization of the polyacrylate resins (H), the comments made above in
25 relation to the polyurethanes (A) apply analogously.

In accordance with the invention it is of advantage to prepare the polyacrylate resins (H) in the presence of

saturated polyurethanes (A), and so this variant is particularly preferred.

Very particularly highly suitable polyacrylate resins
5 (H) are obtained by copolymerizing the olefinically unsaturated monomers (h) described below, of which at least one contains at least one carboxylic acid group and preferably at least one contains at least one hydroxyl group and is substantially free from acid
10 groups.

Examples of suitable monomers (h) are:

Monomers (h1):

15 Hydroxyalkyl esters of acrylic acid, methacrylic acid of another alpha,beta-ethylenically unsaturated carboxylic acid which are derived from an alkylene glycol which is esterified with the acid, or are obtainable by reacting the acid with an alkylene oxide,
20 especially hydroxyalkyl esters of acrylic acid, methacrylic acid or ethacrylic acid in which the hydroxyalkyl group contains up to 20 carbon atoms, such as 2-hydroxyethyl, 2-hydroxypropyl, 3-hydroxypropyl, 3-hydroxybutyl, 4-hydroxybutyl acrylate, methacrylate,
25 ethacrylate or crotonate; 1,4-bis(hydroxy-methyl)cyclohexane, octahydro-4,7-methano-1H-indenedi-methanol or methylpropanediol monoacrylate, mono-methacrylate, monoethacrylate or monocrotonate; or reaction products of cyclic esters, such as epsilon-

caprolactone, for example, and these hydroxyalkyl esters; or olefinically unsaturated alcohols such as allyl alcohol or polyols such as trimethylolpropane monoallyl or diallyl ether or pentaerythritol 5 monoallyl, diallyl or triallyl ether. These monomers (h1) of higher functionality are generally used only in minor amounts. In the context of the present invention, minor amounts of higher-functional monomers here are amounts which do not result in the crosslinking or 10 gelling of the polyacrylate resins. Thus, for example, the proportion of trimethylolpropane monoallyl ether may be from 2 to 10% by weight, based on the overall weight of the monomers (h1) to (h6) used to prepare the polyacrylate resin. In addition, however, it is also 15 possible to add from 2 to 10% by weight, based on the overall weight of the monomers (h) used to prepare the polyacrylate resin, of trimethylolpropane monoallyl ether to the finished polyacrylate resin. Although the olefinically unsaturated polyols (h1) can be used as 20 the sole monomers (h1), it is of advantage in accordance with the invention to use them in combination with further monomers (h1).

Monomers (h2):

25 (Meth)acrylic alkyl or cycloalkyl esters having up to 20 carbon atoms in the alkyl radical, especially methyl, ethyl, propyl, n-butyl, sec-butyl, tert-butyl, hexyl, ethylhexyl, stearyl and lauryl acrylate or methacrylate; cycloaliphatic (meth)acrylic esters,

especially cyclohexyl, isobornyl, dicyclopentadienyl, octahydro-4,7-methano-1H-indenemethanol or tert-butylcyclohexyl (meth)acrylate; (meth)acrylic oxaalkyl esters or oxacycloalkyl esters such as ethyltriglycol (meth)acrylate and methoxyoligoglycol (meth)acrylate having a molecular weight Mn of preferably 550; or other ethoxylated and/or propoxylated hydroxyl-free (meth)acrylic acid derivatives. These may include, in minor amounts, higher-functional (meth)acrylic alkyl or 5 cycloalkyl esters such as ethylene glycol, propylene glycol, diethylene glycol, dipropylene glycol, butylene glycol, 1,5-pentanediol, 1,6-hexanediol, octahydro-4,7-methano-1H-indenedimethanol or cyclohexane-1,2-, -1,3- or -1,4-diol di(meth)acrylate; trimethylolpropane di- 10 or tri(meth)acrylate; or pentaerythritol di-, tri- or tetra(meth)acrylate. In the context of the present invention, minor amounts of higher-functional monomers 15 (h2) here are amounts which do not cause crosslinking or gelling of the polyacrylate resins.

20

Monomers (h3):

Ethylenically unsaturated monomers which carry at least one acid group, preferably a carboxyl group, per molecule, or a mixture of such monomers. As component 25 (h3) it is particularly preferred to use acrylic acid and/or methacrylic acid. It is also possible, however, to use other ethylenically unsaturated carboxylic acids having up to 6 carbon atoms in the molecule. Examples of such acids are ethacrylic acid, crotonic acid,

maleic acid, fumaric acid, and itaconic acid. It is also possible to use ethylenically unsaturated sulfonic or phosphonic acids, and/or their partial esters, as component (h3). Further suitable monomers (h3) include
5 mono(meth)acryloyloxyethyl maleate, succinate, and phthalate.

Monomers (h4):

Vinyl esters of alpha-branched monocarboxylic acids
10 having 5 to 18 carbon atoms in the molecule. The branched monocarboxylic acids may be obtained by reacting formic acid or carbon monoxide and water with olefins in the presence of a liquid, strongly acidic catalyst; the olefins may be cracking products from
15 paraffinic hydrocarbons, such as mineral oil fractions, and may contain both branched and straight-chain acyclic and/or cycloaliphatic olefins. In the reaction of such olefins with formic acid and/or with carbon monoxide and water, a mixture of carboxylic acids is
20 formed in which the carboxyl groups are located predominantly on a quaternary carbon atom. Other olefinic starting materials are, for example, propylene trimer, propylene tetramer, and diisobutylene. Alternatively, the vinyl esters may be prepared in a
25 conventional manner from the acids; for example, by reacting the acid with acetylene. Particular preference - owing to their ready availability - is given to the use of vinyl esters of saturated aliphatic

monocarboxylic acids having 9 to 11 carbon atoms and being branched on the alpha carbon atom.

Monomers (h5):

5 Reaction product of acrylic acid and/or methacrylic acid with the glycidyl ester of an alpha-branched monocarboxylic acid having 5 to 18 carbon atoms per molecule. The reaction of the acrylic or methacrylic acid with the glycidyl ester of a carboxylic acid
10 having a tertiary alpha carbon atom may take place before, during or after the polymerization reaction. As component (h5) it is preferred to use the reaction product of acrylic and/or methacrylic acid with the glycidyl ester of Versatic® acid. This glycidyl ester
15 is obtainable commercially under the name Cardura® E10. For further details, refer to Römpf Lexikon Lacke und Druckfarben, Georg Thieme Verlag, Stuttgart, New York, 1998, pages 605 and 606.

20 **Monomers (h6):**

Ethylenically unsaturated monomers substantially free from acid groups, such as

- olefins such as ethylene, propylene, 1-butene, 1-pentene, 1-hexene, cyclohexene, cyclopentene,
- 25 norbornene, butadiene, isoprene, cyclopentadiene and/or dicyclopentadiene;
- (meth)acrylamides such as (meth)acrylamide, N-methyl-, N,N-dimethyl-, N-ethyl-, N,N-diethyl-, N-propyl-, N,N-dipropyl, N-butyl-, N,N-dibutyl-,

N-cyclohexyl- and/or N,N-cyclohexyl-methyl-(meth)acrylamide;

- monomers containing epoxide groups, such as the glycidyl ester of acrylic acid, methacrylic acid, ethacrylic acid, crotonic acid, maleic acid, fumaric acid and/or itaconic acid;
- vinylaromatic hydrocarbons, such as styrene, alpha-alkylstyrenes, especially alpha-methylstyrene, arylstyrenes, especially diphenylethylene, and/or vinyltoluene;
- nitriles such as acrylonitrile and/or methacrylonitrile;
- vinyl compounds such as vinyl chloride, vinyl fluoride, vinylidene dichloride, vinylidene difluoride; N-vinylpyrrolidone; vinyl ethers such as ethyl vinyl ether, n-propyl vinyl ether, isopropyl vinyl ether, n-butyl vinyl ether, isobutyl vinyl ether and/or vinyl cyclohexyl ether; vinyl esters such as vinyl acetate, vinyl propionate, vinyl butyrate, vinyl pivalate, vinyl esters of Versatic® acids, which are marketed under the brand name VeoVa® by the company Deutsche Shell Chemie (for further details, refer to Römpf Lexikon Lacke und Druckfarben, Georg Thieme Verlag, Stuttgart, New York, 1998, page 598 and also pages 605 and 606), and/or the vinyl ester of 2-methyl-2-ethylheptanoic acid; and/or
- polysiloxane macromonomers having a number-average molecular weight Mn of from 1 000 to 40 000,

preferably from 2 000 to 20 000, with particular preference from 2 500 to 10 000, and in particular from 3 000 to 7 000, and having on average from 0.5 to 2.5, preferably from 0.5 to 1.5,
5 ethylenically unsaturated double bonds per molecule, as are described in DE-A-38 07 571 on pages 5 to 7, in DE-A-37 06 095 in columns 3 to 7, in EP-B-0 358 153 on pages 3 to 6, in US-A-4,754,014 in columns 5 to 9, in
10 DE-A-44 21 823, or in the international patent application WO 92/22615 on page 12, line 18 to page 18, line 10, or acryloxy silane-containing vinyl monomers, preparable by reacting hydroxy-functional silanes with epichlorohydrin and then
15 reacting the reaction product with methacrylic acid and/or hydroxyalkyl esters of (meth)acrylic acid.

In accordance with the invention it is especially
20 advantageous to select the monomers (h) so as to give hydroxyl-containing polyacrylate resins which have an OH number of preferably from 0 to 200, more preferably from 60 to 140, acid numbers of preferably from 20 to 100, more preferably from 25 to 50, glass transition
25 temperatures Tg of preferably from -25 to +80°C, more preferably from -20 to +40°C, and number-average molecular weights of preferably from 1 500 to 30 000, more preferably from 1 500 to 20 000 (determined by gel

permeation chromatography using polystyrene as internal standard).

The glass transition temperature Tg of the polyacrylate
5 resins (H) is determined by the nature and amount of
the monomers (h1) and, where appropriate (h2), (h3),
(h4), (h5) and/or (h6) that are used. The skilled
worker is able to select the monomers with the
assistance of the following formula of Fox, which can
10 be used for approximate calculation of the glass
transition temperatures Tg of (co)polymers, especially
polyacrylate resins:

$$n = x$$

$$\frac{1}{Tg} = \sum_{n=1}^x W_n / T_{gn}$$

$$n = 1$$

15

Tg = glass transition temperature of the
polyacrylate resin

W_n = weight fraction of the nth monomer

T_{gn} = glass transition temperature of the homopolymer
20 of the nth monomer

x = number of different monomers

Viewed in terms of its method, the preparation of the
hydroxyl-containing polyacrylate resins (H) for
25 inventive use has no special features but instead takes
place in accordance with the customary and known

methods of radical polymerization in the presence of at least one polymerization initiator.

Examples of suitable polymerization initiators are
5 initiators which form free radicals, such as tert-butylperoxyethyl hexanoate, benzoyl peroxide, di-tert-amyl peroxide, azobisisobutyronitrile, and tert-butyl perbenzoate. The initiators are used preferably in an amount of from 1 to 25% by weight, with particular
10 preference from 2 to 10% by weight, based on the overall weight of the monomers.

The polymerization is appropriately conducted at a temperature of from 80 to 200°C, preferably from 110 to
15 180°C.

It is preferred to commence the initiator feed a certain time, generally from about 1 to 15 minutes, before the monomers feed. Preference is also given to a process wherein the addition of initiator is commenced at the same time as the addition of the monomers and ended about half an hour after the addition of the monomers has been ended. The initiator is preferably added in a constant amount per unit time. After the end
20 of the addition of initiator, the reaction mixture is held at polymerization temperature until (generally 1.5 hours) all of the monomers used have undergone substantially complete reaction. "Substantially complete reaction" is intended to denote that,

preferably, 100% by weight of the monomers used have been converted but that it is also possible for a small residual monomer content of not more than up to about 0.5% by weight, based on the weight of the reaction mixture, to remain unreacted.

One particularly preferred process for the preparation of the polyacrylate resins (H) is described in DE-A-44 37 535.

10

In this process, in a first step, a polyurethane free from copolymerizable double bonds, especially a polyurethane (A), is introduced in organic solution, after which a mixture of at least one monomer (h1) and 15 at least one monomer (h2) and also, where appropriate, in each case at least one monomer (h4), (h5) and/or (h6) is added, and the mixture is copolymerized. After these monomers (h) have undergone almost complete conversion, in a second process step a mixture of at 20 least one monomer (h3) and also in each case at least one monomer (h1) (h2), (h4), (h5) and/or (h6) is added, after which the resulting reaction mixture is polymerized to completion.

25

In terms of apparatus as well, the preparation of the polyacrylate resins (H) for inventive use has no special features as far as its method is concerned but instead takes place with the aid of the methods, customary and known in the polymers field, of

continuous or batchwise copolymerization under atmospheric or superatmospheric pressure in stirred tanks, autoclaves, tube reactors or Taylor reactors.

5 Examples of suitable copolymerization processes and apparatus are described in the patents DE-A-197 09 465, DE-C-197 09 476, DE-A-28 48 906, DE-A-195 24 182, EP-A-0 554 783, WO 95/27742 or WO 82/02387.

10 Following their preparation, the polyacrylate resins (H) are neutralized with at least one of the above-described suitable neutralizing agents (D), and dispersed in water. The total amount of neutralizing agent (D) used in the aqueous basecoat material of the
15 invention is chosen such that from 1 to 100 equivalents, preferably from 50 to 90 equivalents, of the carboxyl groups of the polyurethanes (A) for inventive use and of the polyacrylate resins (H) are neutralized.

20 The fraction of the polyacrylate resins (H) in the aqueous basecoat materials of the invention can vary widely. It is preferably from 0.1 to 10%, more preferably from 0.5 to 8.0%, with particular preference
25 from 0.6 to 6.0%, with very particular preference from 0.8 to 6.0%, and in particular from 1.0 to 6.0% by weight, based in each case on the overall weight of the respective aqueous basecoat material of the invention.

In yet another preferred embodiment of the aqueous basecoat material of the invention, it comprises at least one water-soluble or -dispersible polyester resin (I).

5

Examples of suitable starting products for preparing the polyester resins (I) are the compounds described above in connection with the preparation of the polyester polyols.

10

Examples of suitable polyester resins (I) and their preparation from said starting products are described in detail in the patents DE-A-40 09 858 or
15 DE-A-44 37 535.

20

The fraction of the polyester resins (I) in the aqueous basecoat materials of the invention can also vary widely. It is preferably from 0.5 to 12%, more preferably from 0.7 to 10%, with particular preference from 0.8 to 9.0%, with very particular preference from 1.0 to 8.0%, and in particular from 1.5 to 7.0% by weight, based in each case on the overall weight of the respective aqueous basecoat material of the invention.

25

In addition to the constituents described above, the aqueous basecoat material of the invention may comprise customary and known additives (J) in effective amounts.

Examples of suitable additives (J) are

- organic and inorganic fillers such as chalk, calcium sulfate, barium sulfate, silicates such as talc or kaolin, silicas, oxides such as aluminum hydroxide or magnesium hydroxide, or organic fillers such as textile fibers, cellulose fibers, polyethylene fibers or wood flour; for further details refer to Römpf Lexikon Lacke und Druckfarben, Georg Thieme Verlag, 1998, pages 250 ff., "Fillers";
- customary and known oligomeric and polymeric binders such as thermally curable, hydroxyl-containing, linear and/or branched and/or block, comb and/or random poly(meth)acrylates or acrylate copolymers, polyesters, alkyds, acrylated polyesters, polylactones, polycarbonates, polyethers, epoxy resin-amine adducts, (meth)acrylate diols, partially hydrolyzed polyvinyl esters, or polyureas;
- customary and known thermally curable reactive diluents such as positionally isomeric diethyloctanedioils or hydroxyl-containing hyper-branched compounds or dendrimers;
- low-boiling and/or high-boiling organic solvents ("long solvents");
- UV absorbers;
- light stabilizers such as HALS compounds, benzotriazoles or oxalanilides;

- free-radical scavengers;
- thermolabile free-radical initiators such as organic peroxides, organic azo compounds or C-C-cleaving initiators such as dialkyl peroxides,
- 5 peroxocarboxylic acids, peroxydicarbonates, peroxide esters, hydroperoxides, ketone peroxides, azo dinitriles or benzpinacol silyl ethers;
- crosslinking catalysts such as dibutyltin dilaurate, lithium decanoate or zinc octoate, or
- 10 amine-blocked organic sulfonic acids;
- devolatilizers, such as diazadicycloundecane;
- slip additives;
- polymerization inhibitors;
- defoamers;
- 15 -
- emulsifiers, especially nonionic emulsifiers such as alkoxylated alkanols and polyols, phenols and alkylphenols or anionic emulsifiers such as alkali metal salts or ammonium salts of alkanecarboxylic acids, alkanesulfonic acids, and sulfo acids of
- 20 alkoxylated alkanols and polyols, phenols and alkylphenols;
- wetting agents such as siloxanes, fluorine compounds, carboxylic monoesters, phosphates, polyacrylic acids and their copolymers, or
- 25 polyurethanes;
- adhesion promoters such as tricyclodecanedi-methanol;
- leveling agents;

- film-forming auxiliaries such as cellulose derivatives;
- transparent fillers based on silica, alumina, titanium dioxide or zirconium oxide; for further details refer to Römpf Lexikon Lacke und Druckfarben, Georg Thieme Verlag, Stuttgart, 1998, pages 250 to 252;
- other rheology control additives, such as those known from patents WO 94/22968, EP-A-0 276 501, EP-A-0 249 201 or WO 97/12945; crosslinked polymeric microparticles, as disclosed for example in EP-A-0 008 127; silicas such as Aerosils; or synthetic polymers containing ionic and/or associative groups such as polyvinyl alcohol, poly(meth)acrylamide, poly(meth)acrylic acid, polyvinylpyrrolidone, styrene-maleic anhydride or ethylene-maleic anhydride copolymers and their derivatives or hydrophobically modified polyacrylates; and/or
- 20 - flame retardants.

Further examples of suitable coatings additives are described in the textbook "Lackadditive" by Johan Bieleman, Wiley-VCH, Weinheim, New York, 1998.

25 The aqueous basecoat materials of the invention preferably have a solids content at spray viscosity of from 5.0 to 60%, more preferably from 5.0 to 50%, with particular preference from 10 to 45%, with very

particular preference from 13 to 40%, and in particular from 13 to 35% by weight, based in each case on the overall weight of the respective aqueous basecoat material of the invention.

5

The preparation of the aqueous basecoat material of the invention has no special features but instead takes place in a customary and known manner by mixing of the above-described constituents in appropriate mixing

10 equipment such as stirred tanks, dissolvers or extruders in accordance with the techniques suitable for preparing the respective aqueous basecoat materials.

15 The aqueous basecoat material of the invention is used to produce the coatings of the invention, especially multicoat paint systems, on primed or unprimed substrates.

20 Suitable substrates are all surfaces for coating which are not damaged by curing of the coatings present thereon using heat. Suitable substrates comprise, for example, the moldings, films and fibers of the invention, metals, plastics, wood, ceramic, stone,
25 textile, fiber assemblies, leather, glass, glass fibers, glass wool and rock wool, mineral-bound and resin-bound building materials, such as plasterboard panels and cement slabs or roof shingles, and composites of these materials. The coating system of

- the invention, accordingly, is also suitable for applications outside of automotive finishing, particularly automobile finishing. In such applications it is suitable particularly for the coating of
- 5 furniture and for industrial coating, including coil coating, container coating, and the impregnation or coating of electrical components. In the context of industrial coatings it is suitable for coating virtually all parts for private or industrial use such
- 10 as radiators, domestic appliances, small metal parts such as nuts and bolts, hubcaps, wheel rims, packaging, or electrical components such as motor windings or transformer windings.
- 15 In the case of electrically conductive substrates it is possible to use primers, which are prepared in a customary and known manner from electrocoat materials. Both anodic and cathodic electrocoat materials are suitable for this purpose, but especially cathodics.
- 20 With the multicoat paint system of the invention it is also possible to coat primed or unprimed plastics such as, for example, ABS, AMMA, ASA, CA, CAB, EP, UF, CF, MF, MPF, PF, PAN, PA, PE, HDPE, LDPE, LLDPE, UHMWPE,
- 25 PET, PMMA, PP, PS, SB, PUR, PVC, RF, SAN, PBT, PPE, POM, PUR-RIM, SMC, BMC, PP-EPDM and UP (abbreviations to DIN 7728T1). The plastics to be coated can of course also be polymer blends, modified plastics or fiber-reinforced plastics. It is also possible to employ the

plastics that are commonly used in vehicle construction, especially motor vehicle construction.

In the case of unfunctionalized and/or apolar substrate surfaces, these may be subjected prior to coating in a known manner to a pretreatment, such as with a plasma or by flaming, or may be provided with a water-based primer.

10 The multicoat paint systems of the invention may be produced in a variety of ways.

A first preferred variant of the process of the invention comprises the following steps:

15 (I) preparing a basecoat film by applying the aqueous basecoat material of the invention to the substrate,
(II) drying the basecoat film,
(III) preparing a clearcoat film by applying a clearcoat material to the basecoat film, and
20 (IV) jointly curing the basecoat film and the clearcoat film, to give the basecoat and the clearcoat (wet-on-wet technique).

25 This variant offers particular advantages especially in the context of the coating of plastics, and is therefore employed with particular preference in that utility.

A second preferred variant of the process of the invention comprises the following steps:

(I) preparing a surfacer film by applying a surfacer to the substrate,

5 (II) curing the surfacer film, to give the surfacer coat,

(III) preparing a basecoat film by applying the aqueous basecoat material of the invention to the surfacer coat,

10 (IV) drying the basecoat film,

(V) preparing a clearcoat film by applying a clearcoat material to the basecoat film, and

(VI) jointly curing the basecoat film and the clearcoat film, to give the basecoat and the clearcoat (wet-on-wet technique).

A third preferred variant of the process of the invention comprises the following steps:

(I) preparing a surfacer film by applying a surfacer to the substrate,

20 (II) drying the surfacer film,

(III) preparing a basecoat film by applying the aqueous basecoat material of the invention to the surfacer film,

25 (IV) drying the basecoat film,

(V) preparing a clearcoat film by applying a clearcoat material to the basecoat film, and

(VI) jointly curing the surfacer film, the basecoat film and the clearcoat film, to give the basecoat and the clearcoat (extended wet-on-wet technique).

5 The two last-mentioned variants offer particular advantages especially in the context of the coating of automobile bodies and are therefore employed with very particular preference in that utility.

10 It is found here to be a further particular advantage of the aqueous basecoat material of the invention and of the process of the invention that all customary and known clearcoat materials can be combined with the aqueous basecoat film of the invention in the context
15 of the processes of the invention.

Examples of suitable known one-component (1K), two-component (2K) or multicomponent (3K, 4K) clearcoat materials are known from the patents DE-A-42 04 518,

20 US-A-5,474,811, US-A-5,356,669, US-A-5,605,965, WO 94/10211, WO 94/10212, WO 94/10213, EP-A-0 594 068, EP-A-0 594 071, EP-A-0 594 142, EP-A 0 604 992, WO 94/22969, EP-A-0 596 460 or WO 92/22615.

25 One-component (1K) clearcoat materials are known to contain hydroxyl-containing binders and crosslinking agents such as blocked polyisocyanates, tris-(alkoxycarbonylamino)triazines and/or amino resins. In another variant they include as binders polymers

containing pendant carbamate and/or allophanate groups and carbamate- and/or allophanate-modified amino resin crosslinking agents (cf. US-A-5,474,811, US-A-5,356,669, US-A-5,605,965, WO 94/10211, 5 WO 94/10212, WO 94/10213, EP-A-0 594 068, EP-A-0 594 071 or EP-A-0 594 142).

Two-component (2K) or multicomponent (3K, 4K) clearcoat materials include as essential constituents, as is known, hydroxyl-containing binders and polyisocyanate crosslinking agents, which are stored separately until they are used.

Examples of suitable powder clearcoat materials are known, for example, from the German patent DE-A-42 22 194 or from the BASF Lacke + Farbe AG product information bulletin "Pulverlacke", 1990.

Powder clearcoat materials include as essential constituents, as is known, binders containing epoxide groups, and polycarboxylic acid crosslinking agents.

Examples of suitable powder slurry clearcoat materials are known, for example, from the US patent US-A-4,268,542 and from the German patent applications DE-A-195 18 392.4 and DE-A-196 13 547, or are described in the German patent application DE-A-198 14 471.7, unpublished at the priority date of the present specification.

Powder slurry clearcoat materials, as is known, comprise powder clearcoat materials in dispersion in an aqueous medium.

5

UV-curable clearcoat materials are disclosed, for example, by the patents EP-A-0 540 884, EP-A-0 568 967 or US-A-4,675,234.

10 Furthermore, the clearcoats may additionally be coated further with at least one other clearcoat, an example being an organically modified ceramic layer, thereby making it possible to improve significantly the scratch resistance of the multicoat paint system of the
15 invention.

Accordingly, the multicoat paint systems of the invention can differ in their structure.

20 In a first preferred variant of the multicoat paint system of the invention, the following coats lie above one another in the stated order:

(1) a surfacer coat which absorbs mechanical energy,

25

(2) the color and/or effect basecoat, and

(3) a clearcoat.

In the second preferred variant of the multicoat paint system of the invention, the following coats lie above one another in the stated order:

5 (1) the color and/or effect basecoat, and

(2) a clearcoat. This preferred variant is used in particular in the coating of plastics.

10 The aqueous basecoat material of the invention may be applied by all customary application methods, such as spraying, knifecoating, brushing, flowcoating, dipping, impregnating, trickling, or rolling, for example. The substrates to be coated may itself be at rest, with the
15 application equipment or unit being moved. Alternatively, the substrate to be coated, especially a coil, may be moved, with the application unit being at rest relative to the substrate or being moved appropriately.

20

Preference is given to the use of spray application methods, such as compressed-air spraying, airless spraying, high-speed rotation, electrostatic spray application (ESTA), alone or in conjunction with hot
25 spray application such as hot-air spraying, for example. Application may be conducted at temperatures of max. 70 to 80°C, so that suitable application viscosities are achieved without the short-term thermal stress being accompanied by any change in or damage to

the aqueous basecoat material or its overspray, which may be intended for reprocessing. For instance, hot spraying may be configured such that the aqueous basecoat material is heated in the spray nozzle for
5 only a very short time, or is heated just a short way upstream of the spray nozzle.

The spray booth used for the application may be operated, for example, with an optionally temperature-
10 controllable circulation, which is operated with an appropriate absorption medium for the overspray, an example being the aqueous basecoat material itself.

In general, the surfacer film, basecoat film and
15 clearcoat film are applied in a wet film thickness such that they cure to give coats having the coat thicknesses which are necessary and advantageous for their functions. In the case of the surfacer coat, this coat thickness is from 10 to 150, preferably from 15 to
20 120, with particular preference from 10 to 100, and in particular from 10 to 90 μm ; in the case of the basecoat it is from 5 to 50, preferably from 5 to 40, with particular preference from 5 to 30, and in particular from 10 to 25 μm ; and in the case of the
25 clearcoats it is from 10 to 100, preferably from 15 to 80, with particular preference from 20 to 70, and in particular from 25 to 60 μm .

The surfacer film, basecoat film and clearcoat film are cured thermally.

Full curing may take place after a certain rest time.

- 5 Its duration may be from 30 s to 2 h, preferably from 1 min to 1 h, and in particular from 1 min to 45 min. The rest time serves, for example, for the coating films to flow and undergo devolatilization, or for the evaporation of volatile constituents such as solvents.
- 10 The rest time may be assisted and/or shortened by the application of elevated temperatures of up to 90°C and/or by a reduced atmospheric humidity < 10 g water/kg air, especially < 5 g water/kg air, provided this does not entail any damage or change to the
- 15 coating films, such as premature complete crosslinking, for instance.

The thermal curing has no special features in terms of its method but instead takes place in accordance with

- 20 the customary and known methods, such as heating in a convection oven or irradiation with IR lamps. This thermal curing may also take place in stages. Advantageously, thermal curing takes place at a temperature of from 50 to 100°C, with particular
- 25 preference from 80 to 100°C, and in particular from 90 to 100°C, for a time of from 1 min up to 2 h, with particular preference from 2 min up to 1 h, and in particular from 3 min to 30 min. If substrates of high heat resistance are used, thermal crosslinking may also

be conducted at temperatures above 100°C. In this case it is generally advisable not to exceed temperatures of 180°C, preferably 160°C, and in particular 155°C.

- 5 In the case of clearcoats, depending on the clearcoat material used, curing may also take place with actinic radiation or both thermally and with actinic radiation (dual cure). Suitable actinic radiation includes electromagnetic radiation such as near infrared light
10 (NIR), visible light, UV radiation or X-rays, and/or corpuscular radiation such as electron beams.

The multicoat paint systems of the invention exhibit an outstanding profile of properties which is very well balanced in terms of mechanics, optics, corrosion resistance, and adhesion. Thus the multicoat paint systems of the invention possess the high optical quality and intercoat adhesion required by the market and do not give rise to any problems such as deficient
20 surfacer-coat condensation resistance, cracking (mud-cracking) in the basecoats or leveling defects or surface structures in the clearcoats.

In particular, the multicoat paint system of the
25 invention exhibit an outstanding metallic effect, an excellent D.O.I. (distinctiveness of the reflected image), and an outstanding surface smoothness. They are weathering-stable, resistant to chemicals and bird

droppings, are scratch resistant, and exhibits very good reflow behavior.

Not least, however, it proves to be a very special
5 advantage that the use of the aqueous basecoat materials of the invention in the production of the multicoat paint systems of the invention results in only barely visible optical defects, if any.

10 Accordingly, the substrates coated with them have particular advantages such as a prolonged service life, a better esthetic impression on the viewer, and improved technological usefulness, so making them particularly economically attractive.

15

Examples and comparative experiments

Preparation example 1

20 **The preparation of an aqueous polyurethane dispersion (A)**

716.6 parts by weight of a condensation product (number-average molecular weight: 1410) of 1.81 mol of
25 a polymeric fatty acid (dimer content at least 98% by weight, trimer content not more than 2.0% by weight, monomer content traces at most), 0.82 mol of isophthalic acid, 0.61 mol of hexanediol and 0.61 mol of neopentyl glycol, 61 parts by weight of

dimethylolpropionic acid, 10.6 parts by weight of neopentyl glycol, 365 parts by weight of methyl ethyl ketone and 308.3 parts by weight of m-tetramethylxylylidene diisocyanate were heated at 80
5 degrees Celsius under nitrogen and with stirring in a suitable reaction vessel. Reaction was continued to an isocyanate content of 1.1% by weight, based on the overall amount of reaction mixture. Subsequently 52.6 parts by weight of trimethylolpropane were added, after
10 which the resulting reaction mixture was stirred at 80 degrees Celsius until free isocyanate groups were no longer detectable. Thereafter, slowly, 33 parts by weight of dimethylethanolamine, 255 parts by weight of butyl glycol and, subsequently, 2153 parts by weight of
15 deionized water were stirred in. The methyl ethyl ketone was distilled off under reduced pressure. This gave a fine dispersion whose pH was adjusted to 7.4 using dimethylethanolamine and whose nonvolatiles fraction was adjusted to 31% by weight using deionized
20 water.

Preparation example 2

The preparation of an aqueous polyester resin solution

25 (I)

A reactor equipped with stirrer, thermometer and packed column was charged with 729 parts by weight of neopentyl glycol, 768 parts by weight of hexanediol,

462 parts by weight of hexahydrophthalic anhydride and
1710 parts by weight of a polymeric fatty acid (dimer
content at least 98% by weight, trimer content not more
than 2% by weight, monomer content traces at most), and
5 this initial charge was melted. The resulting melt was
heated with stirring in such a way that the overhead
column temperature did not exceed 100 degrees Celsius.
Esterification was carried out at 220 degrees Celsius
maximum until an acid number of 9 was reached. After
10 the mixture had cooled to 180 degrees Celsius, 768
parts by weight of trimellitic anhydride were added and
esterification was continued until an acid number of 32
was reached. Thereafter, the reaction mixture was
cooled to 120 degrees Celsius and diluted with 1392
15 parts by weight of butyl glycol. After it had cooled to
90 degrees Celsius, 158 parts by weight of
dimethylethanolamine were stirred in slowly, followed
by 1150 parts by weight of deionized water. The
resulting polyester resin solution was adjusted to a pH
20 of 7.6 using dimethylethanolamine and to a nonvolatiles
fraction of 60% by weight using deionized water.

Preparation example 3

25 **The preparation of a polyacrylate resin (H)**

500 parts by weight of a condensation product (number-
average molecular weight: 1423) of 1.0 mol of a
polymeric fatty acid (dimer content at least 98% by

- weight, dimer content not more than 2.0% by weight, monomer content traces at most), 1.5 mol of isophthalic acid, 1.6 mol of neopentyl glycol and 1.7 mol of hexanediol, 31.2 parts by weight of neopentyl glycol,
5 185 parts by weight of methyl ethyl ketone, 201.7 parts by weight of m-tetramethylxylylidene diisocyanate and 0.7 part by weight of dibutyltin dilaurate were heated at 80 degrees Celsius under nitrogen and with stirring. The reaction was continued to an isocyanate content of
10 1.3% by weight, based on the overall amount of reaction mixture. Thereafter, 30 parts by weight of diethanolamine were added and the resulting reaction mixture was stirred at 80 degrees Celsius until isocyanate groups were no longer detectable.
15 Subsequently, 466 parts by weight of butyl glycol were stirred in, after which the methyl ethyl ketone was distilled off under reduced pressure. The resulting polyurethane solution was then adjusted to a nonvolatiles fraction of 60% by weight using butyl
20 glycol.

A steel vessel equipped with monomer feed, initiator feed, thermometer, oil heating and reflux condenser was charged with 28.44 parts by weight of butyl glycol and
25 24.24 parts by weight of the abovementioned polyurethane solution, and this initial charge was heated to 110 degrees Celsius. Then a solution of 5.1 parts by weight of t-butyl perethylhexanoate in 6.0 parts by weight of butyl glycol was added uniformly at

a rate such that the addition was over after 5.5 hours. The beginning of the addition of initiator was also accompanied by commencement of the addition of a mixture of 18.36 parts by weight of n-butyl
5 methacrylate, 17 parts by weight of methyl methacrylate, 17 parts by weight of lauryl methacrylate, 17.34 parts by weight of hydroxypropyl acrylate and 12.75 parts by weight of styrene. The monomer mixture was added uniformly at a rate such that
10 the addition was over in five hours. After all of the initiator solution had been added, the reaction mixture was held at 110 degrees Celsius for a further hour.

Subsequently, a solution of 1.17 parts by weight of
15 t-butyl perethylhexanoate in 3.5 parts by weight of butyl glycol was added uniformly at a rate such that the addition was over after 1.5 hours. The beginning of the addition of the initiator solution was also accompanied by the commencement of addition of a mixture of 5.85 parts by weight of acrylic acid and
20 4.65 parts by weight of n-butyl methacrylate, 2.94 parts by weight of methyl methacrylate, 5.90 parts by weight of lauryl methacrylate, 1.25 parts by weight of hydroxypropyl acrylate and 2.94 parts by weight of
25 styrene. The monomer mixture was added uniformly at a rate such that the addition was over within one hour. Thereafter, the temperature of the reaction mixture was held at 110 degrees Celsius for 1.5 hours more. The resulting resin solution was concentrated by

distillation under reduced pressure to a solids content
of 80% by weight and at this temperature was
neutralized with dimethylethanolamine over the course
of 30 minutes to a degree of neutralization of 80%. The
5 resin solution was cooled to 60 degrees Celsius, after
which the heating was shut off. Then water was added
slowly until the solids content of the dispersion was
40% by weight. The dispersion had an acid number of
36.7 mg KOH/g and a pH of 7.6.

10

Examples 1 and 2 and comparative experiments C1 and C2

The preparation of inventive (examples 1 and 2) and
noninventive (comparative experiments C1 and C2)
15 aqueous basecoat materials

The preparations of the inventive aqueous basecoat
materials of examples 1 and 2 and of the noninventive
aqueous basecoat material of comparative experiment C2
20 were conducted in analogy to the preparation
instructions indicated below for the noninventive
aqueous basecoat material of comparative experiment C1.
Table 1 gives an overview of the amounts of the
starting products employed.

25

For comparative experiment C1, the thickener 1 (paste
of a synthetic sodium magnesium phyllosilicate from
Laporte; 3% in water) was introduced initially. Added
to this with stirring were deionized water, thickener 3

(3% aqueous solution of a polyacrylic acid from Allied Colloids; trade name: Viscalex), the polyurethane dispersion (A) from preparation example 1, the polyester resin solution (I) from preparation example 5 2, the solution of the polyacrylate (H) from preparation example 3, butyl glycol, a commercial melamine resin in butanol (Maprenal® VMF3924), a silica filler paste (12 parts by weight of Syloid® ED-3 from Grace; 30 parts by weight of the abovementioend 10 polyester resin solution (I)), a neutralizing agent (D) (dimethylethanolamine, 10% in water), a solution of a commercial defoamer based on alkynediol (50% in butyl glycol; from Air Products), deionized water, a pigment paste 1 (63 parts by weight of polyurethane dispersion, 15 31 parts by weight of titanium rutile Tayca® MT500HD from Tayca), a pigment paste 2 (64 parts by weight of polyurethane dispersion and 32 parts by weight of titanium dioxide UVL 530 from Kemira).

20 Prepared separately from this was an aluminum pigment slurry composed of 5.0 parts by weight of a commercial aluminum paste (Alu-Stapa-Hydrolux® 2192 from Eckart), 6.0 parts by weight of solvent and 2.4 parts by weight of the polyester resin solution (I). This aluminum 25 pigment slurry was stirred into the mixture described above.

The resultant mixture was then neutralized to a pH of about 8.0 using the neutralizing solution (D) and was

adjusted using deionized water to a viscosity of 65 mPas under a shear of 1291 s⁻¹.

- For comparative experiment C2, comparative experiment
5 C1 was repeated but using the thickener 2 instead of the thickener 1. Said thickener 2 is a 3% solution of a synthetic sodium magnesium phyllosilicate from Südchemie.
- 10 For example 1, comparative experiment C1 was repeated but using the thickener 4 instead of the thickener 3. Said thickener 4 is a mixture of 50% by weight of a polyurethane-based associative thickener (G) (Nopco® DSX1550 from Henkel) and 50% by weight of butyl glycol.
- 15 For example 2, example 1 was repeated but using, instead of butyl glycol, dipropylene glycol monoalkyl ether for the preparation of the aluminum pigment slurry.
- 20 The aqueous basecoat materials of comparative experiment C2 and of examples 1 and 2 were also adjusted using neutralizing solution (D) to a pH of about 8.0 and using deionized water to a viscosity of 25 65 mPas under a shear of 1291 s⁻¹.

Table 1 gives an overview of the amounts of the starting products and of the composition of the aqueous basecoat materials.

Table 1: The composition of the inventive (examples 1 and 2) and of the noninventive (comparative experiments C1 and C2) aqueous basecoat materials

5

Constituents	Comparative experiments:		Examples:	
	C1	C2	1	2
Thickener 1	29	-	29	29
Thickener 2	-	29	-	-
Deionized water	9.0	9.0	9.0	9.0
Thickener 3	5.5	5.5	-	-
Polyurethane dispersion (A)	21.5	21.5	21.5	21.5
Polyester solution (I)	2.0	2.0	2.0	2.0
Polyacrylate resin solution (H)	5.0	5.0	5.0	5.0
Butyl glycol	2.0	2.0	2.0	2.0
Melamine resin (B)	5.0	5.0	5.0	5.0
Filler paste	1.2	1.2	1.2	1.2
Neutralizing agent (D)	0.4	0.4	0.4	0.4
Defoamer	2.0	2.0	2.0	2.0
Thickener 4	-	-	1.6	1.6
Deionized water	4.0	4.0	4.0	4.0
Pigment paste 1	4.3	4.3	4.3	4.3
Pigment paste 2	0.8	0.8	0.8	0.8
Alu-Stapa-Hydrolux®	5.0	5.0	5.0	5.0
Butyl glycol	6.0	6.0	6.0	-
Dipropylene glycol methyl ether	-	-	-	6.0
Polyester solution (I)	2.4	2.4	2.4	2.4

The viscosity behavior of the inventive (examples 1 and 2) and of the noninventive (comparative experiments C1 and C2) aqueous basecoat materials was measured under 5 different shear rates. The results can be found in tab. 2.

Table 2: The viscosity behavior of the inventive (examples 1 and 2) and of the noninventive 10 (comparative experiments C1 and C2) aqueous basecoat materials

Shear rate (s^{-1})	Viscosity (mPas):			
	Comparative experiments:		Examples:	
	C1	C2	1	2
34 (outgoing curve)	360	220	355	350
1000	73	73	73	73
34 (return curve)	250	180	245	235

The values from table 2 demonstrate that the aqueous 15 basecoat materials of examples 1 and 2 and of comparative experiments C1 and C2 had a comparable (comparative experiment C2 on the one hand and examples 1 and 2 and comparative experiment C1 on the other) or identical (examples 1 and 2 and comparative example C1) 20 viscosity behavior.

Examples 3 and 4 and comparative experiments C3 and C4

Testing of the inventive (examples 3 and 4) and of the noninventive (comparative experiments C3 and C4)
5 aqueous basecoat materials for their action against the formation of optical defects

For example 3, the inventive aqueous basecoat material of example 1 was used.

10

For example 4, the inventive aqueous basecoat material of example 2 was used.

15

For comparative experiment C3, the noninventive aqueous basecoat material of comparative experiment C1 was used.

20

For comparative experiment C4, the noninventive aqueous basecoat material of comparative experiment C2 was used.

The test panels of examples 3 and 4 and of comparative experiments C3 and C4 were prepared in accordance with the following general instructions:

25

Steel panels coated cathodically with a commercial electrocoat material (electrocoat with a thickness of 18-22 µm) were first coated with a commercial filler from BASF Coatings AG, using a cup-type gun, and baked.

This gave a surfacer with a thickness of from 35 to 40 μm . Atop the surfacer there were then applied, in the same way, the inventive and the noninventive aqueous basecoat materials (cf. table 1), which were 5 predried at 80°C for 10 minutes. Following the cooling of the panels, a coat of commercial two-component high-solids clearcoat material from DuPont was applied in each case, predried at room temperature for 10 minutes, and then crosslinked together with the basecoats at 10 140°C for 45 minutes. This gave basecoats with a thickness of 15 μm and clearcoats with a thickness of 44 μm .

15 1. Light spots

In the wet sanding of defects in an original finish or in a surfacer, water droplets remain on the parts of the body that are to be overcoated with basecoat and clearcoat. These water droplets are made up of water, 20 abrasion dust from clearcoat and/or surfacer, and residues of the abrasive paper. Following the drying of these water droplets and subsequent coating with the basecoat and clearcoat, the sites of the dried-out water are visible as light (-colored) spots.

25 In order to simulate this effect, abrasion water droplets (deionized water + abrasion dust from surfacer and clearcoat + abrasive paper from Plochmann, softened in this composition) of various sizes were applied to

the respective test panels, dried at room temperature and coated with the respective basecoat materials employed for the original finish and described in table 1 and with the two-component high-solids clearcoat 5 material from DuPont. The resulting color differences in the coatings (light spots) were assessed visually. The results obtained can be found in table 3.

2. Polishing spots

10 Polishing sites were produced on the test panels using a red/white felt pad (Buffing Pad from 3M, No. 049-5765) and the gray polishing paste from 3M (No. 029-2778). In the subsequent coating with the 15 basecoat materials described in table 1 and the abovementioned clearcoat material, these polishing sites were clearly visible on account of their different color. The respective difference in color was determined visually. The results obtained can be found 20 in table 3.

3. Abrasion spots

Using an abrasive paper from Plochmann with a grade of 25 2000, both a round and a cross-shaped abrasion site were applied to the test panels. These abrasion sites were coated with the basecoat materials described in table 1 and with the abovementioned clearcoat material.

The respective color difference was determined visually. The results obtained can be found in table 3.

The color difference, i.e., the obviousness of the defects, was scored as follows:

Rating meaning

1	defects not visible
10	defects very slightly visible
3	defects slightly visible
4	defects markedly visible
5	defects very markedly visible

15 Table 3: The effect of the inventive (examples 3 and 4) and of the noninventive (comparative experiments C3 and C4) aqueous basecoat materials on the formation of optical defects

Optical defects	Comparative experiments:		Examples:	
	C3	C4	3	4
Light spots	4	5	2	2
Polishing spots	4	5	4	2-3
Abrasion spots	3-4	4	2-3	2

The results demonstrate that by means of the aqueous basecoat materials of the invention the formation of optical defects can be effectively suppressed.

An additional improvement results from the use of dipropylene glycol monoalkyl ether. The comparison of the results in table 3 with the aqueous basecoat material viscosity behavior shown in the overview in table 2 underlines the fact that said behavior, surprisingly, is not responsible for the advantageous effect of the aqueous basecoat materials of the invention.

Claims

1. Use of polyurethane-based associative thickeners and/or dipropylene glycol monoalkyl ethers for 5 suppressing optical defects in multicoat color and/or effect paint systems or their refinishes.

2. The use as claimed in claim 1, characterized in that the multicoat color and/or effect paint 10 system and/or its refinish comprise or comprises at least one basecoat and at least one clearcoat.

3. The use as claimed in claim 2, characterized in that the basecoat is produced from an aqueous 15 basecoat material.

4. The use as claimed in claim 3, characterized in that the aqueous basecoat material is 20 polyurethane-based.

5. The use as claimed in any of claims 2 to 4, characterized in that the polyurethane-based 25 associative thickener and/or dipropylene glycol monoalkyl ether is or are present in the basecoat material or materials.

6. The use as claimed in any of claims 2 to 5, characterized in that the polyurethane-based

associative thickener and dipropylene glycol monoalkyl ether are present in the basecoat material or materials.

5 7. The use as claimed in any of claims 1 to 5, characterized in that the optical defects comprise

10 - light-colored spots, introduced by dried-up liquid residues and/or abrasion dust residues, which remain after the abrading of defects in a multicoat color and/or effect paint system or in a surfacer that is to be coated with a multicoat color and/or effect paint system; and

15 - polishing spots, induced by the polishing of defects in a multicoat color and/or effect paint system which is performed for the purpose of refinishing the multicoat color and/or effect paint system.

20 8. The use as claimed in any of claims 3 to 7, characterized in that the aqueous basecoat material comprises

25 (A) at least one water-soluble or -dispersible polyurethane,

(B) at least one crosslinking agent,

- (C) at least one color and/or effect pigment,
 - (D) at least one neutralizing agent,
- 5 (E) at least one inorganic thickener, and
- (F) at least one dipropylene glycol monoalkyl ether.
- 10 9. The use as claimed in claim 8, characterized in that the aqueous basecoat material comprises
- (G) at least one polyurethane-based associative thickener.
- 15 10. The use as claimed in any of claims 3 to 7, characterized in that the aqueous basecoat material comprises
- (A) at least one water-soluble or -dispersible polyurethane,
 - (B) at least one crosslinking agent,
- 20 (C) at least one color and/or effect pigment,
- (D) at least one neutralizing agent,
 - (E) at least one inorganic thickener,

(G) at least one polyurethane-based associative thickener,

5 (H) if desired, at least one water-soluble or -dispersible polyacrylate resin prepared in the presence of a water-soluble or -dispersible polyurethane, and

10 (I) at least one water-soluble or -dispersible polyester resin.

11. The use as claimed in claim 10, characterized in that the aqueous basecoat material comprises

15 (F) at least one dipropylene glycol monoalkyl ether.

12. The use as claimed in any of claims 8, 9 and 11,
20 characterized in that the aqueous basecoat material, based on its overall weight, contains from 0.5 to 11% by weight of dipropylene glycol monoalkyl ethers (F).

25 13. The use as claimed in any of claims 9 to 12, characterized in that the aqueous basecoat material, based on its overall weight, contains from 0.1 to 4% by weight of at least one polyurethane-based associative thickener (G).

14. The use as claimed in any of claims 3 to 13,
characterized in that the aqueous basecoat
material comprises

5

- (J) at least one additive.

15. An aqueous basecoat material comprising

10 (A) at least one water-soluble or -dispersible
polyurethane,

- (B) at least one crosslinking agent,

15 (C) at least one color and/or effect pigment,

- (D) at least one neutralizing agent,

- (E) at least one inorganic thickener, and

20 (F) at least one dipropylene glycol monoalkyl
ether.

16. The aqueous basecoat material as claimed in claim

25 15, characterized in that it comprises

- (G) at least one polyurethane-based associative
thickener.

17. An aqueous basecoat material comprising

(A) at least one water-soluble or -dispersible polyurethane,

5

(B) at least one crosslinking agent,

(C) at least one color and/or effect pigment,

10 (D) at least one neutralizing agent,

(E) at least one inorganic thickener,

(G) at least one polyurethane-based associative
15 thickener,

(H) if desired, at least one water-soluble or -
dispersible polyacrylate resin prepared in
the presence of a water-soluble or -
dispersible polyurethane, and

20 (I) at least one water-soluble or -dispersible
polyester resin.

25 18. The aqueous basecoat material as claimed in claim
17, characterized in that it comprises

(F) at least one dipropylene glycol monoalkyl
ether.

19. The aqueous basecoat material as claimed in any of

claims 15, 16 and 18, characterized in that it,
based on its overall weight, contains from 0.5 to
5 11% by weight of dipropylene glycol monoalkyl
ethers.

20. The aqueous basecoat material as claimed in any of

claims 16 to 19, characterized in that it, based
10 on its overall weight, contains from 0.1 to 4% by
weight of at least one polyurethane-based
associative thickener.

21. The aqueous basecoat material as claimed in any of

15 claims 15 to 20, characterized in that it
comprises

(J) at least one additive.

20 22. The use of the aqueous basecoat material as

claimed in any of claims 15 to 21 for the original
coating of motor vehicle bodies, industrial
components, including electrical components,
coils, and packaging, plastics or furniture, or
25 for the refinish of multicoat color and/or effect
paint systems present thereon.

23. Multicoat color and/or effect paint systems

comprising at least one basecoat producible from

an aqueous basecoat material as claimed in any of
claims 15 to 21.

ABSTRACT

Use of associative thickeners based on polyurethane and/or dipropylene glycol monoalkyl ethers for suppressing defects in color and/or effect multicoat paint systems, or their touch-up paints.

DECLARATION FOR UTILITY OR DESIGN PATENT APPLICATION (37 CFR 1.63)		Attorney Docket No. IN - 5562	
<input checked="" type="checkbox"/> Declaration or <input type="checkbox"/> Declaration		First Named Inventor Ines MANGELS et al.	
Submitted with initial Filing (surcharge (37 CFR 1.16 (e))		COMPLETE IF KNOWN	
Submitted after initial Filing		Application Number	
		Filing Date	Monday, March 11, 2002
		Group Art Unit	
		Examiner Name	

As below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

USE OF ASSOCIATIVE THICKENERS BASED ON POLYURETHANE AND/OR DIPROPYLENE GLYCOL MONOALKYL ETHERS IN COLOURED AND/OR DECORATIVE EFFECT MULTI-LAYERED LACQUERS

(Title of the Invention)

The specification of which:

is attached hereto

Was filed on 31. August 2000 as United States Application or PCT International Application Number PCT/EP00/08503
and was amended on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment specifically referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56, including for continuation-in-part applications, material information which became available between the filing date of the prior application and the national or PCT international filing date of the continuation-in-part application.

I hereby claim foreign priority benefits under 35, U.S.C § 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate or 365 (a) of any PCT international application(s) designating at least one country other than the United States of America listed below and have also identified below, by checking the box, any foreign application(s) for patent or inventor's certificate, or any PCT international application(s) having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Foreign Filing Date DATE/MONTH/YEAR	Priority Not Claimed	Certified Copy Attached? Yes	Certified Copy Attached? No
199 45 574.0	GERMANY	23. September 1999	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Additional foreign application number are listed on a supplemental priority data sheet PTO/SB/02B attached hereto:

I hereby claim the benefit under 35 U.S.C 119(e) of any United States provisional application(s) listed below:

APPLICATION NUMBER(S)	FILING DATE

Additional provisional application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto

DECLARATION —**Utility or Design Patent Application****POWER OF ATTORNEY**

I hereby appoint the practitioner(s) associated with the Customer Number provided below to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith

<input type="checkbox"/> Customer Number or Bar Code Label	 26922 <small>PATENT TRADEMARK OFFICE</small>	or <input type="checkbox"/> Correspondence address below
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County	United States of America	Telephone	(248) 948-2019	Fax	(248) 948-2093
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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Name OF SOLE OR FIRST INVENTOR:		<input type="checkbox"/> A petition has been filed for this unsigned inventor					
Given Name (first and middle [if any])		Family Name or Surname					
Ines		MANGELS					
Inventor's Signature		Date					
Signature		31.01.2002					
Residence: City	48165 Münster	State	Germany	Country	Federal Republic of Germany	Citizenship	German
Mailing Address		Pinienweg 21					
City	48165 Münster	State	Germany	Zip			

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<input type="checkbox"/>	Customer Number or Bar Code Label		or	<input type="checkbox"/> Correspondence address below
26922 PATENT-TMADKMA OFFICE				

County	United States of America	Telephone	(248) 948-2019	Fax	(248) 948-2093
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Name OF SECOND INVENTOR: <i>J. Sten</i>				A petition has been filed for this unsigned inventor			
Given Name (first and middle if any)		<u>Karl-Heinz</u>		Family Name or Surname		<u>JOSTEN</u>	
Inventor's Signature		<i>J. Sten</i>		Date		<u>05.02.2002</u>	
Residence: City	97209	State	<u>Germany</u> <i>(Signature)</i>	Country	Federal Republic of Germany	Citizenship	German
Mailing Address	Brusseler Str. 8						
City	97209 Veitshöchheim	State	<u>Germany</u>	Zip			

DECLARATION

Utility or Design Patent Application

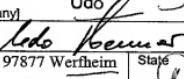
POWER OF ATTORNEY

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<input type="checkbox"/>	Customer Number or Bar Code Label		or	Correspondence address below
26922 PATENT-TRADEMARK OFFICE				

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Name of Additional Joint Inventor, if any:		<input type="checkbox"/> A petition has been filed for this unsigned inventor					
Given Name (first and middle if any)	Udo	Family Name or Surname					
Inventor's Signature		Date					
Residence: City	97877 Wertheim	State	Germany	Country	Federal Republic of Germany	Citizenship	German
Mailing Address	Am Mühlacker 3						
City	97877 Wertheim	State	Germany	Zip			

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<input checked="" type="checkbox"/>	Customer Number or Bar Code Label		or	<input type="checkbox"/>	Correspondence address below
26922 PATENT-TMADMARK OFFICE					

County	United States of America	Telephone	(248) 948-2019	Fax	(248) 948-2093
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Name of Additional Joint Inventor, if any:			<input type="checkbox"/> A petition has been filed for this unsigned inventor		
Given Name (first and middle [if any])		Egon	<i>4-10</i>		
Inventor's Signature		<i>C. Wegner</i>			
Residence: City	97209 Veitshöchheim	State	Germany <i>Weg</i>		
Mailing Address	Martinstr. 8				
City	97209 Veitshöchheim	State	Germany	Zip	

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Name of Additional Joint Inventor, if any:				<input type="checkbox"/> A petition has been filed for this unsigned inventor
Given Name (first and middle if any)	Ekkehard	Signature	Family Name or Surname	SAPPER
Inventor's Signature	Date			17. Februar 2002
Residence: City	97222 Rimpar	State	Country	Federal Republic of Germany
Mailing Address	Austr. 55		Citizenship	German
City	97222 Rimpar	State	Germany	Zip

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<input checked="" type="checkbox"/>	Customer Number or Bar Code Label		or	<input type="checkbox"/>	Correspondence address below
26922 PATENT-TRADEMARK OFFICE					

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Name of Additional Joint Inventor, If any:				<input type="checkbox"/>	A petition has been filed for this unsigned inventor			
Given Name (first and middle if any)		Nicola		<input type="checkbox"/>	Family Name or Surname			
Inventor's Signature		<i>Nicola Zarse</i>		Date	<i>04.07.02</i>			
Residence: City	48165 Münster	State	Germany	Country	Federal Republic of Germany	Citizenship	German	
Mailing Address		Kardinalstr.16						
City	48165 Münster	State	Germany	Zip				